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A biochemical and genetic discovery pipeline identifies PLC δ 4b as a non-receptor activator of heterotrimeric G proteins

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ABSTRACT

Recent evidence has revealed that heterotrimeric G proteins can be activated by cytoplasmic proteins that share an evolutionarily conserved sequence called the G α -binding-and-activating (GBA) motif. This mechanism alternative to canonical activation by GPCRs plays important roles in cell function and its dysregulation is linked to disease such as cancer. Here we describe a discovery pipeline that uses biochemical and genetic approaches to validate GBA candidates identified by sequence similarity. First, putative GBA motifs discovered in bioinformatics searches were synthesized on peptide arrays and probed in batch for G α i3 binding. Then, cDNAs encoding proteins with G α i3-binding sequences were expressed in a genetically-modified yeast strain that reports mammalian G protein activity in the absence of GPCRs. The resulting GBA motif candidates were characterized by comparison of their biochemical, structural, and signaling properties with those of all previously described GBA motifs in mammals (GIV/Girdin, DAPLE, Calnuc, and NUCB2). We found that the phospholipase C δ 4 (PLC δ 4) GBA motif binds G proteins with high affinity, has guanine-nucleotide exchange factor activity *in vitro*, and activates G-protein signaling in cells, as indicated by bioluminescence resonance energy

transfer (BRET)-based biosensors of G-protein activity. Interestingly, the PLC δ 4 isoform b, which lacks the domains required for PLC activity, bound and activated G proteins more efficiently than the full-length isoform a, suggesting that PLC δ 4b functions as a G-protein regulator rather than as a PLC. In summary, we have identified PLC δ 4 as a non-receptor activator of G proteins, and established an experimental pipeline to discover and characterize GBA motif-containing proteins.

Heterotrimeric G proteins are gatekeepers of signal transduction that regulate fundamental aspects of cell biology across the full spectrum of the eukaryota domain, from unicellular fungi to multicellular mammals (1,2). They work as molecular switches that are turned on or off depending on their guanine nucleotide binding status. Thus, the biological activity of heterotrimeric G proteins is determined by their nucleotide handling properties (such as GDP/GTP exchange and GTP hydrolysis), which control the amplitude and duration of downstream signaling. At the same time, nucleotide binding and hydrolytic activity is modulated by a large network of regulators (2). The best characterized components of this network are membrane receptors of the GPCR¹ superfamily. In the canonical paradigm, ligand-activated GPCRs act as Guanine-nucleotide Exchange Factors (GEFs)

that promote the exchange of GDP for GTP in the $G\alpha$ subunit of a resting $G\alpha\beta\gamma$ heterotrimer. As a consequence of GTP binding, $G\alpha$ and $G\beta\gamma$ dissociate from each other and subsequently activate downstream effectors that propagate signaling intracellularly. In turn, signaling is terminated upon hydrolysis of GTP by $G\alpha$, which returns it to the GDP-bound form that reassociates with $G\beta\gamma$. These series of reactions constitute the so-called canonical G protein cycle (1).

In addition to the components described above as core elements of the canonical regulatory cycle, G proteins interact with a wide range of “accessory proteins” (3). These are cytoplasmic factors that can be classified based on their biochemical activity towards G proteins. The best characterized of these accessory proteins are members of the Regulators of G protein Signaling (RGS) family, which bind to active G proteins and have GTPase Activating Protein (GAP) activity (4,5), thereby facilitating the termination of G protein signaling via acceleration of GTP hydrolysis. This family of proteins shares a signature domain of ~120 aa called the RGS box, which confers the GAP activity (6-9). Another well characterized family of accessory proteins is defined by a signature sequence of 30-40 aa called the GoLoco motif, which confers Guanine nucleotide Dissociation Inhibitor (GDI) activity (10-12). Members of this family also bind to $G\alpha$ subunits, but do so when they are in the inactive, GDP-bound form to prevent nucleotide exchange and subsequent activation (13-15). Both RGS and GoLoco proteins have been shown to play crucial roles in cell biology and human disease (12,16,17).

A third group of accessory proteins can be broadly defined based on their biochemical activity as GEFs. These are cytoplasmic factors that mimic the action of GPCRs, i.e., they promote the exchange of GDP for GTP on $G\alpha$ subunits. However, in contrast to RGS and GoLoco proteins, the initial discovery of non-receptor GEFs was not accompanied by the identification of a shared domain or motif responsible for the biochemical activity, as this group was constituted by unrelated proteins and the domains responsible for the G protein regulatory activity were not mapped (18-22). The exception to this was GIV (also known as Girdin), for which a short sequence of ~30 aa named the $G\alpha$ -Binding-and-Activating

(GBA) motif was identified (23,24). Subsequently, it has been shown that the GBA motif is present in evolutionarily unrelated proteins and defines a mechanism of G protein activation by non-receptor proteins already existing in invertebrates (25-27). Thus, the GBA motif represents the defining structural element for a subfamily of non-receptor G protein activators with GEF activity (28). In mammals, this group currently consists of four proteins (GIV, DAPLE, Calnuc and NUCB2), which bind and activate $G\alpha$ subunits of the Gi subfamily (i.e., $G\alpha i1$, $G\alpha i2$ and $G\alpha i3$). The structural basis for the binding and activation of Gai proteins by the GBA motif *in vitro* has been studied in detail (29-31), and it has also been shown that the GBA motif activates G proteins in cells using approaches that monitor the formation of Gai-GTP (cAMP or conformation-specific antibodies) or free $G\beta\gamma$ (live-cell biosensors or PI3K-Akt signaling) (23,26,32-36). Such signaling mechanisms have a profound impact on cellular processes such as cell migration, mitosis, polarity or autophagy (24,26,37-39), which underlie the linkage between GBA protein dysregulation and human disorders such as cancer, hepatic fibrosis, renal dysfunction, insulin resistance or pathologic vascularization (24,40).

Despite these advances in characterizing the structural basis, cell signaling and disease linkage properties of GBA proteins, a fundamental question that remains unanswered is how prevalent this mechanism of G protein regulation is. For this, a critical first step is to identify which proteins, and how many, contain a functional GBA motif. In this regard, the amino acid sequence of the GBA motif of proteins identified to date is known, but this information is insufficient to systematically identify other GBA motifs based solely on sequence similarity. Although sequence similarity has been successfully used to identify members of other families of G protein regulators (e.g., RGS GAPs), the short length of the GBA motif and the limited number of validated GBA sequences limits the power of bioinformatics searches. For these reasons, bioinformatics searches for GBA motifs yield a large number of false positives. To overcome this limitation, here we describe a pipeline of biochemical and genetic assays that can be used to validate and characterize GBA candidates identified by sequence similarity to

currently known GBA proteins. Using peptide arrays, functional assays in genetically-engineered yeast and other biochemical assays, we identify PLC84 as a novel GBA motif-containing non-receptor GEF and subsequently characterize its signaling properties by systematic comparison with all the other GBA motifs described to date.

RESULTS

Identification of putative GBA motifs by bioinformatics and peptide arrays- We envisioned a screening strategy to discover GBA motif-containing G protein activators (**Fig. 1A, B**). We started by using bioinformatics to identify putative GBA motifs by sequence similarity, followed by a high-throughput peptide array assay to identify Gai3 binders as initial hits. The subsequent validation pipeline involved genetic assays in yeast and further biochemical characterization. The underlying logic of this approach is that previous evidence indicates that Gai3 binding is an absolute requirement for GBA-mediated G protein activation, and that the extent of binding and GEF activity correlate tightly (23,25,26,29,31). For this, we used the consensus sequence of the 7 amino acids of the highly conserved core of the GBA motifs identified to date (23,25,36,41-43) to search for similar sequences using ScanSite3 (44). This program uses a Position Specific Scoring Matrix (PSSM), which in our case was defined as [VLIM]-[T]-[VLIM]-x-[DE]-F-[VLIM] based on the consensus GBA motif ψ -T- ψ -x-D/E-F- ψ , where ψ is an aliphatic residue and x is any residue (**Fig. 1A**). Approximately 200 hits were obtained using this PSSM to search the human proteome based on the most stringent cut-off setting of ScanSite3 (0.2 score). Proteins that were annotated in UniProt (45) as secreted/ extracellular or as transmembrane proteins in which the predicted GBA sequence was either in the membrane spanning region or not facing the cytosol were eliminated. The 88 candidate GBA motifs that resulted from this curation were synthesized as 25mer peptides (in which the invariable F of the GBA motif occupied the center position) immobilized on cellulose and probed in batch with purified His-Gai3 (**Fig. 1C**). This set of peptides included sequences derived from DAPLE and Calnuc, two previously validated GBA motifs that served as internal controls. We found that DAPLE was the strongest

binder in the peptide array (**Fig. 1C**), which is consistent with previous observations showing that DAPLE has the highest affinity for Gai3 of all the GBA motifs described to date ($K_d \sim 0.1$ - $0.4 \mu\text{M}$) (26,29). On the other hand, Calnuc binding was weaker (**Fig. 1C**), which is also consistent with the K_d previously reported for its interaction with Gai3 ($K_d \sim 1$ - $4 \mu\text{M}$) (25). We reasoned that peptides binding Gai3 at least as much as the Calnuc GBA peptide might represent physiologically relevant interactors because Calnuc has been previously shown to interact directly with Gai3 in a cellular context (25,46). Based on this criterion, 29 GBA candidates were selected for further investigation.

Validation of GBA candidate hits as G protein activators in yeast- While binding of Gai3 to immobilized GBA peptides suggests that these sequences are capable of G protein binding in isolation, this might not be the case when the sequence is placed in the context of a folded protein and/ or might be insufficient for activation of G protein signaling. To start validating the GBA candidates, we used a genetic approach with a previously described genetic screen approach in yeast (47,48) (**Fig. 2A**). Briefly, we expressed the cDNA of GBA candidates in a genetically-engineered yeast strain that lacks GPCRs and with the endogenous yeast G α protein Gpa1 replaced by human Gai3. In this “humanized” yeast system, only an exogenous G protein activator can trigger a signaling pathway that is normally activated as a pheromone response that leads to an increase in transcriptional activation of the *FUS1* gene (**Fig. 2A**). The cDNAs of GBA candidates were cloned as full-length proteins based on naturally occurring isoforms annotated in UniProt or as truncations corresponding to domains known or predicted to fold as independent units (see details in **Fig. S1**). These constructs were expressed in the genetically-engineered yeast strain described above and G protein activation was assessed by the activation of a *FUS1* promoter β -galactosidase reporter (**Fig. 2B**). As an internal control we used Ric-8A, a protein that displays GEF activity toward Gai proteins *in vitro* (19) and that we have previously validated as a G protein activator in this yeast-based assay (27). From the 29 candidates identified as Gai3 binders in the peptide array (**Fig. 1**), 8 were toxic in yeast or excluded for

other reasons (**Fig. S1**). From the remaining 21 candidate genes expressed in yeast, PLCD4 (which encodes for the phospholipase PLC δ 4) stood out from the rest by inducing *FUS1* activation ~80-fold over basal levels. Another 6 candidates (NOC2L, K1045, KCIP1, KALIRIN, ALKB4 and MAPK2) induced *FUS1* activation of much lower magnitude (~3-8-fold) than PLCD4 but still comparable to the positive control Ric-8A (**Fig. 2B**). As a secondary validation of these results, we measured Fus3 phosphorylation in lysates of the same yeast cells used in the reporter assay. Because Fus3 phosphorylation is an event upstream of *FUS1* transcriptional activation in the signaling cascade triggered by G proteins (**Fig. 2A**), we reasoned that GBA candidates leading to increased β -galactosidase activity should also have increased Fus3 phosphorylation. We found that this was the case except for KCIP1 (**Fig. 2C**), which failed to increase Fus3 phosphorylation despite leading to *FUS1* activation based on β -galactosidase activity measurements. We concluded that KCIP1's effect on β -galactosidase activity is likely due to a G protein-independent mechanism, while the GBA candidates PLCD4, NOC2L, K1045, KALIRIN, ALKB4 and MAPK2 might be G protein activators.

Validation of GBA candidate hits as direct G protein binders- As an independent validation of GBA candidates, we also investigated if they bound directly to G proteins. As mentioned above, it has been previously shown that binding of GBA proteins to Gai3 correlates tightly with their ability to activate G protein signaling (23,25,26,29,31), so we established binding as a selection criterion to identify *bona fide* GBA motif-containing G protein activators. For this, we cloned the exact same sequences of the 29 GBA candidate genes used for the yeast assays described above as GST-fusions for expression in *E. coli* (**Fig. 1B**). We could purify 15 of them in sufficient quantity and quality for protein-protein binding experiments. This set included one protein that could not be tested in the yeast-based assays due to toxicity (ABLIM3). RCN2 and TF3B were toxic in yeast and could not be purified from *E. coli*, but subsequent pulldown experiments with lysates of mammalian cells expressing these candidates ruled out binding to Gai3 (**Fig. S2**).

Binding of purified His-Gai3 to the 15 GST-fused GBA candidates purified from *E. coli* was tested in pulldown assays using permissive conditions (i.e., relatively high concentrations of proteins) to facilitate the identification of binders. As a positive control, we used GST-Calnuc. As indicated above, Calnuc is a validated GBA protein that binds with relatively weak affinity compared to other GBA motif-containing proteins (25,46), so we considered positive for binding only those GBA candidates that displayed binding similar to that detected with Calnuc. Based on this criterion, KCIP1, KALIRIN, ALKB4 and MAPK2 did not bind to His-Gai3 (**Fig. 3**). Thus, it is likely that the activity of these candidates in the yeast-based system (**Fig. 2B, C**) is exerted via G protein-independent mechanisms, and we excluded them from further characterization as possible GBA motif containing G protein activators. On the other hand, the other three GBA candidates that were active in the yeast assays (**Fig. 2B, C**), i.e., PLCD4, NOC2L and K1045, did bind directly to His-Gai3 (**Fig. 3**).

PLCD4 and NOC2L, but not K1045, activate G protein signaling in yeast via their GBA motif-

Next, we investigated if the GBA motif identified in PLCD4, NOC2L and K1045 in our bioinformatics searches (**Fig. 1**) is responsible for the ability of these proteins to activate G protein signaling in yeast (**Fig. 2**). For this, we mutated the conserved phenylalanine (F) in the GBA motif (**Fig. 1A**) of each one of these candidates to alanine (A) and assessed the effect of these mutants on β -galactosidase *FUS1* reporter activity and Fus3 phosphorylation in the “humanized” yeast strain described above (**Fig. 2A**). It has been previously shown that equivalent “FA” mutations in any of the previously validated GBA motif-containing proteins (GIV, DAPLE, Calnuc, NUCB2, GBAS-1) abolish G protein binding and GEF activity almost completely (23,25-27,29). Consistent with these previous observations, we found that the FA mutants of PLCD4 and NOC2L failed to recapitulate the activation exerted by their wild-type (WT) counterparts in either β -galactosidase *FUS1* reporter activity assays (**Fig. 4A**) or Fus3 phosphorylation immunoblots (**Fig. 4B**). On the other hand, the equivalent mutation in K1045 did not have an effect in the same assays (**Fig. 4A, B**), suggesting that the GBA motif of

this protein is not responsible for its G protein regulatory activity. Taken together, these results indicate that PLCD4 and NOC2L activate G protein signaling via their GBA motifs.

PLCD4 and NOC2L bind preferentially to GDP-bound Gai3 over the GTP-bound form- Much like other proteins with GEF activity, all previously characterized GBA proteins (GIV, DAPLE, Calnuc, NUCB2, GBAS-1) bind preferentially to GDP-bound, inactive Gai3 over GTP-bound Gai3 (23,25-27,29). Next, we investigated if this was the case for PLCD4 and NOC2L. Purified His-Gai3 was preloaded with GDP, GDP/AlF₄⁻ (which mimics the GTP-bound transition state (49,50)) or GTPγS (which is a non-hydrolyzable GTP analog) by incubation at 30 °C for 3 hours, and used as the soluble ligand in pulldowns with resin-immobilized GST-PLCD4. We found that PLCD4 binding to Gai3 loaded with either of the two GTP mimetics (GDP/AlF₄⁻ or GTPγS) was much weaker than to Gai3-GDP (**Fig. 5A**), suggesting that PLCD4 has G protein state-dependent binding properties analogous to those previously described for other proteins with GBA motifs. The same nucleotide preincubation conditions of 3 hours at 30 °C diminished the already weak binding of NOC2L to Gai3-GDP to undetectable levels, which precluded any comparison with GTP mimetics. To overcome this limitation, we omitted the GTPγS condition and performed the preloading step at 4 °C for 20-30 minutes, which is sufficient to load the G protein with AlF₄⁻. Using these modified preincubation conditions, we found that NOC2L binds preferentially to inactive (Gai3-GDP) vs. active (Gai3-GDP/AlF₄⁻) (**Fig. 5B**). Interestingly, K1045 showed the opposite pattern of binding, i.e., it bound preferentially to GTP mimetic-bound Gai3 over Gai3-GDP (*not shown*), which along with the lack of effect of the FA mutant in G protein activity assays in yeast (**Fig. 4**) supports that this protein does not behave like other GBA proteins. Studies on the significance of the K1045-Gai3 interaction are being pursued out of the scope of this manuscript.

PLCD4 GBA motif, but not NOC2L GBA motif, binds with high affinity to Gai3- Next, we set out to characterize the binding properties of the GBA motifs of PLCD4 and NOC2L. We have recently

shown that a 31mer peptide corresponding to the GBA motif of GIV is sufficient to recapitulate the high affinity binding ($K_d \sim 0.5\text{--}0.6 \mu\text{M}$) of the protein (30). Using NMR spectroscopy, we also found that the G protein binding contacts of a GIV GBA peptide are the same as for a larger 200 aa fragment of GIV (1660-1870) (29) that is necessary and sufficient to activate G proteins (31,42). Consistently, the NMR spectroscopy studies also showed that the conformational changes in Gai3 associated with G protein activation by GIV were the same for the GBA peptide and the larger 200 aa construct (36). Thus, peptides corresponding to the GBA motif can recapitulate well the G protein binding properties of the proteins they are embedded in. In order to quantify the Gai3 binding affinity of the GBA motifs of PLCD4 and NOC2L compared to previously described GBA motifs in mammalian proteins (GIV, DAPLE, Calnuc and NUCB2), we synthesized fluorescently-labeled peptides and used them in fluorescence polarization (FP) assays to calculate the K_d . Following the design of the GIV GBA motif peptide, we generated analogous 31-mer peptides corresponding to the GBA motifs of DAPLE, Calnuc, NUCB2, PLCD4 and NOC2L (**Fig. 6A**), which would be expected to yield an increase in FP upon binding to Gai3 (~40 kDa). Consistent with previously published data, the GIV GBA motif bound Gai3 with a $K_d \sim 0.6 \mu\text{M}$ (**Fig. 6B**). DAPLE, Calnuc and NUCB2 GBA motifs also bound to Gai3 with K_d 's of ~0.4 μM , ~1 μM and ~3 μM , respectively, in this assay (**Fig. 6B**), which are in agreement with the binding affinities of larger fragments of the corresponding proteins using other assays (25,26). These results further support that isolated GBA motifs recapitulate well the G protein binding properties of the proteins they are embedded in. As for PLCD4 and NOC2L, we found that the former binds with high affinity to Gai3 while the latter does not (**Fig. 6B**). PLCD4 GBA motif binds Gai3 with a $K_d \sim 0.25 \mu\text{M}$ (**Fig. 6B**), which is even lower than that of the GBA motif of GIV or of DAPLE. Since GIV and DAPLE have been shown to regulate G protein signaling in cells via GBA-dependent binding (23,26,31,36,37,51), this suggests that the GBA motif of PLCD4 might also be bioactive. On the other hand, NOC2L binding was so weak that the FP signal did not reach saturation even at the highest concentration of G

protein tested (32 μ M) (**Fig. 6B**), thereby precluding K_d calculation. We estimate that the K_d for the NOC2L GBA: Gai3 interaction is at least one order of magnitude larger than the K_d for the weakest GBA motif (NUCB2). This low affinity of the isolated GBA peptide likely reflects the G protein binding properties of the NOC2L protein because G protein activation in yeast assays and direct Gai3 binding were also weak (compared to PLCD4) in previous experiments using a larger domain of NOC2L (**Fig. 2, Fig. 3**). Although we cannot rule out that the weak affinity interaction of NOC2L with Gai3 is biologically relevant, we focused our subsequent efforts on characterizing the GBA motif of PLCD4.

Identification of critical residues involved in forming the Gai3:PLCD4 interface- To start characterizing the structural properties of the interface formed between PLCD4 and Gai3, we generated a homology-based model of the complex. For this, we used the previously described crystal structure of a GBA-like synthetic peptide called KB-752 bound to Gai1 as a template (41). This model resolves a 13 amino acid stretch of PLCD4 containing the core of the GBA motif. As expected, this model predicts that the GBA motif of PLCD4 binds to Gai3 similar to other GBA motifs, i.e., it docks onto the cleft formed by the Switch II (SwII) region and α 3 helix of Gai3 (**Fig. 7A**). Consistent with this mode of binding, we found that myristoylation of the N-terminus of Gai, which is far from the SwII/ α 3 pocket in crystal structures, does not affect binding to the GBA motif of PLCD4 (**Fig. S3**). To validate more extensively the predicted conservation between the mode of binding of PLCD4 GBA motif and that of other GBA motifs, we tested its binding to a battery of 18 Gai3 mutants previously used to characterize the binding of GIV and DAPLE GBA motifs (29). This set includes mutants within or near the SwII/ α 3 region previously shown to inhibit, enhance or have no effect on binding to the GBA motif of GIV or DAPLE (29). We reasoned that this systematic approach would allow us to define a “fingerprint” of PLCD4 GBA motif binding properties based on the effect of different mutants and that this fingerprint could be compared to that of previously characterized GBA motifs (29). For this, we performed FP experiments with a

fluorescently labeled PLCD4 GBA peptide to quantify the K_d for each mutant and then calculated the fold change in affinity compared to Gai3 WT (**Fig. 7B**). This dataset for PLCD4 GBA motif was compared to previously obtained data (29) for GIV GBA motif and DAPLE GBA motif using the same experimental conditions. As shown in the heat map and correlation plots in **Fig. 7B**, there was good agreement between the effects of Gai3 mutants on the binding of PLCD4 GBA motif and on GIV or DAPLE GBA motif. The Pearson’s correlation coefficient was 0.89 for PLCD4/ GIV and 0.95 for PLCD4/ DAPLE. In general, mutants that diminished affinity for GIV or DAPLE also had diminished affinity for PLCD4, and mutants that increased affinity, like S252A, also did so for all the GBA motifs (**Fig. 7B**). A notable exception was the effect of the W258F mutation. While this mutation *diminishes* the affinity for GIV (~3.5 fold) and DAPLE (~2-fold), it *increases* affinity for PLCD4 (~2 fold). Although our homology model does not include the predicted contacts made by W258 with the GBA motif (**Fig. 7A**), it is reasonable to think that the difference in binding upon W258 mutation arises from the divergence of the sequence in the C-terminal region (i.e., downstream of the conserved F) of the GBA motifs of PLCD4, GIV and DAPLE (**Fig. 6A**). Overall, these results indicate that, barring some nuanced differences, the GBA motif of PLCD4 physically engages G proteins in a manner similar to previously characterized GBA motifs.

Consistent with the homology model (**Fig. 7A**), some of the most marked defects in GBA motif binding were observed upon mutation of the aromatic residues W211 and F215 in Gai3 (**Fig. 7B**), which are predicted to form extensive hydrophobic contacts with PLCD4 GBA motif. More specifically, residues L224, F227 and L228 of PLCD4 are located on one side of the α -helical region of the GBA motif that docks onto the Gai3 hydrophobic groove dominated by W211 and F215 (**Fig. 7A, right**). Again, this is a feature conserved with previously described GBA motifs, as all of them have hydrophobic residues in these positions (**Fig. 1A**) and mutation of the invariable phenylalanine almost completely abolishes binding for GIV, DAPLE, Calnuc and NUCB2 (23,25,26). When we tested if this was also the

case for PLCD4, we found that mutation of F227 to alanine (“FA”) caused a marked but incomplete decrease in Gai3 binding compared to PLCD4 GBA WT (**Fig. 7C**). This incomplete reduction in binding is also in agreement with the incomplete ablation of activity in yeast-based assays using the same mutant (**Fig. 4A, B**). We reasoned that PLCD4 binding to Gai3 might be stabilized by contacts established by the other conserved hydrophobic residues in the GBA motif (i.e., L224, L228), so we tested if multi-position mutants F227A/L228A (“FALA”) and L224A/F227A/L228A (“LAFALA”) were more efficient in disrupting the interaction. We found that this is the case because both FALA and LAFALA displayed undetectable binding to Gai3 (**Fig. 7C**). These results not only indicate that this GBA:G protein interface has properties slightly different from those described for other GBA motifs, as suggested by the Gai3 mutants profiling results (**Fig. 7A**), but also identify the double mutant F227A/L228A as a useful tool to completely blunt the physical coupling of PLCD4 GBA motif to Gai3.

PLCD4 GBA motif activates Gai3 in vitro-

Having identified a powerful tool to assess the specificity of PLCD4 GBA motif action (i.e., the FALA mutant), we set out to determine if this motif had GEF activity like previously characterized GBA proteins and related synthetic peptides (23,25-27,41,43). We have previously validated that steady-state GTPase assays accurately reflect the effect of the GBA proteins GIV, DAPLE, Calnuc and NUCB2 on Gai3 nucleotide exchange, providing results that are indistinguishable from those obtained from GTP γ S binding experiments done in parallel (25,26,29,42). This is explained well by the much faster rate of GTP hydrolysis *versus* nucleotide exchange (2 orders of magnitude for Gai) (52), which makes GDP/GTP exchange the rate limiting step under steady-state GTPase conditions. Using this assay, we found that increasing amounts of GST-fused PLCD4 GBA motif increased the rate of nucleotide exchange on purified His-Gai3 more than 2-fold (**Fig. 7D**), which is comparable to levels previously observed for other regulators of the same family (23,25-27,41,43). This GEF activity was specific because parallel reactions with the G protein binding-deficient PLCD4

FALA mutant revealed no activity (**Fig. 7D**). When analogous experiments were performed with Gai3-G $\beta\gamma$ heterotrimers instead of with free Gai3, we found that PLCD4 WT, but not the FALA mutant, also lead to increased G protein activity, although somewhat less efficiently (~1.7 fold for the trimer vs. ~2 fold for free Gai3) (**Fig. 7D**). Similar results were obtained for His-GIV-CT (aa 1660-1870, containing the GBA motif), in that it also increased activity with Gai3-G $\beta\gamma$ heterotrimers but somewhat less efficiently than with free Gai3 (~2 fold for the trimer (**Fig. 7D**) vs. ~2.5-3 fold for Gai3 (23,31,42). The lower GEF efficacy on heterotrimers compared to monomeric Gai is consistent with previously reported competition between GBA motifs and G $\beta\gamma$ for binding to Gai (23,26). These results indicate that the GBA motif of PLCD4 has GEF activity *in vitro*.

PLCD4 GBA motif activates G protein signaling in mammalian cells-

Next, we investigated if the GBA motif of PLCD4 works as an activator of G proteins in cells. For this, we took advantage of a recently developed experimental system used to demonstrate that membrane recruitment of GIV is sufficient to trigger G protein activation in cells (36,53). The underlying principle of this system is that Gai subunits are constitutively attached to cellular membranes while GIV is predominantly in the cytosol, such that only upon recruitment of GIV (or its GBA motif in isolation) to membranes can it activate G proteins (36,53). Experimentally, membrane recruitment is achieved via chemically-induced dimerization (CID) of FKBP and FRB domains with rapamycin (54) (**Fig. 8A, left**). FRB is fused to the first 11 aa of Lyn (Lyn11-FRB), which targets it to the plasma membrane (55,56), whereas FKBP is fused to a GBA sequence (FKBP-GBA) and remains in the cytosol. Thus, rapamycin induces the translocation of FKBP-GBA to the plasma membrane, where Lyn11-FRB localizes. The effect of this translocation on G protein activation is monitored in real time by bioluminescence resonance energy transfer (BRET) using a genetically-encoded biosensor system. In this system, dissociation of G $\alpha\beta\gamma$ heterotrimers upon activation leads to the association of free Venus-G $\beta\gamma$ (BRET acceptor) with the C-terminal domain of its effector GRK3 fused to an enhanced luciferase (Nluc, BRET

donor), which leads to an increase in BRET (57,58) (**Fig. 8A**, right). Using this experimental system, the amplitude of G protein activation upon GIV GBA translocation is only slightly lower than that obtained upon activation of a Gi-coupled GPCR like the muscarinic M4 receptor with saturating concentrations of ligand (36,53). To assess the G protein regulatory activity of PLCD4 and systematically compare it to other GBA motifs of GIV, DAPLE, Calnuc and NUCB2 as well as of PLCD4 (**Fig. 8B**), and expressed them in HEK293T cells along with Lyn11-FRB and the components of the BRET biosensor. As in our previous report (36), addition of rapamycin induced a rapid increase in BRET in cells expressing FKBP-GBA GIV WT but not FKBP-GBA GIV FA (**Fig. 8C**), indicating GBA-dependent G protein activation upon membrane translocation. Similarly, all the other GBA motifs, including PLCD4, led to an increase in BRET upon rapamycin stimulation that was not recapitulated by the corresponding FA mutants (**Fig. 8C**). However, the amplitude of the BRET responses was different depending on the specific GBA motif used (**Fig. 8C**). In fact, the order of potency of the GBA motifs in this readout matched very well with their affinity for Gai3 determined by FP using the exact same 31-mer GBA sequences (**Fig. 6A**), with PLCD4 as the most potent activator and highest affinity binder. Also consistent with the relative affinity of different GBA motifs for Gai3, we found that the corresponding GBA sequence of NOC2L only induced a marginal increase of BRET in this assay, which was almost negligible in comparison with PLCD4 (**Fig. S4**). One difference between the FKBP-GBA motif of PLCD4 and the other proteins is that the FA mutation failed to completely abolish the BRET response (**Fig. 8C**). This is not surprising given that this mutant still binds weakly to Gai3 (**Fig. 7C**) and moderately activates G protein signaling in yeast (**Fig. 4A, B**). When PLCD4 mutants “FALA” and “LAFALA”, which completely disrupt Gai3 binding and GEF activity *in vitro* (**Fig. 7C, D**), were compared side by side with FKBP-PLCD4 GBA WT and the FA mutant, we observed no measurable G protein activation upon rapamycin stimulation as determined by BRET (**Fig. S5**). These results

confirm the specificity of the GBA-mediated activation of G proteins by PLCD4.

To further substantiate the conclusion that PLCD4 GBA activates G protein signaling in cells, we investigated whether it regulates a signaling event downstream of G protein activation (rather than measuring activation at the level of the G protein as described above). The canonical effectors of G proteins of the Gi family are adenylyl cyclases (AC), the enzymes responsible for the synthesis of cAMP. It is well established that upon Gi activation, GTP-bound Gai subunits bind AC to negatively modulate its activity (59). We used the same rapamycin-induced translocation system as above to trigger the action of FKBP-fused GBA motifs of PLCD4 and GIV (as a control) on G proteins (**Fig. 8D**). The output detection system was a BRET-based biosensor (Nluc-EPAC-VV) that monitors intracellular cAMP levels in real time (57,60) (**Fig. 8D**). Briefly, HEK293T cells were stimulated with forskolin to induce the direct activation of AC and subsequently treated with rapamycin to translocate the GBA motifs to the plasma membrane. While rapamycin had no effect on the forskolin-induced cAMP levels in control cells not expressing any FKBP-GBA fusion, it led to a substantial decrease of cAMP (~25%) in cells expressing FKBP-GBA GIV or FKBP-GBA PLCD4 (**Fig. 8D, E, F**). The response observed for FKBP-GBA PLCD4 was abolished when the same experiment was repeated using the Gai-binding deficient FALA mutant (**Fig. S6**). Although the amplitude of the GBA motif-mediated responses was not as marked as the one observed upon maximal activation of a Gi-coupled GPCR like the α_2A adrenergic receptor, which led to a ~40-45% reduction in cAMP (**Fig. 8F**), these findings are consistent with the activation of Gi-dependent signaling by the GBA motifs of both GIV and PLCD4. Taken together, these results demonstrate that the GBA motif of PLCD4 can activate G protein signaling in cells.

Gai3 binds more strongly to PLC δ 4b than to PLC δ 4a- While the results above strongly indicate that the GBA motif of PLCD4 leads to robust G protein activation, we set out to characterize if this was the case for the GBA motif in its native context of PLCD4 proteins. Based on the curated

annotations for human PLCD4 in UniProt, this gene encodes for two different isoforms of PLC δ 4, i.e., PLC δ 4a (61) and the shorter isoform PLC δ 4b (62) (**Fig. 9A**). PLC δ 4a is considered the canonical isoform of PLC δ 4 and contains all of the domains characteristic of this family (in order from N to C terminus): PH domain that can bind phosphoinositides, EF-hands, catalytic TIM barrel (with X- and Y-boxes) responsible for the phospholipase activity, and C-terminal C2 domain (**Fig. 9A**). Interestingly, PLC δ 4 contains 2 canonical EF-hands (although EF-hands in PLC δ or PLC β isoforms do not bind Ca²⁺ (63-65)) followed by an EF-hand-like domain (203-237 amino acids) of uncharacterized function. The GBA motif of PLCD4 characterized in this work overlaps with this EF-hand-like domain, suggesting that G protein regulation might be its actual function. It is also interesting that the PLC δ 4b isoform contains this EF-hand-like/ GBA sequence but is truncated before the catalytic TIM barrel, which makes it a protein without PLC activity but potentially with G protein regulatory activity. This idea is supported by our results in yeast-based experiments (**Fig. 4A, B**), which showed GBA-dependent G protein activation by a PLCD4 construct that closely resembles PLC δ 4b (i.e., corresponding to amino acids 1-260) (**Fig. 9A**).

The finding of a GBA sequence that overlaps with an EF-hand fold is not novel, as Calnuc and NUCB2 also share this property (25). Moreover, results from our bioinformatics searches and peptide arrays (**Fig. 1**) showed an enrichment for GBA motifs located in EF-hands, i.e., 10 out of the 29 GBA hits in the peptide array correlated with EF-hands. These findings are explained well by the overlap of the consensus GBA motif sequence Ψ -T- Ψ -x-D/E-F- Ψ with the C-terminal segment of the consensus EF-hand sequence D-x-D-x-D-G- Ψ -D/S/T-x-x-D/E-F (underlines indicate the overlap). However, our data also demonstrates that G protein binding (and regulation) is not a widespread function of EF-hands because only one (i.e., PLCD4) of the 10 candidate GBA motifs turned out to have G protein regulatory activity. Instead, G protein regulatory activity seems to correlate with atypical EF-hand sequences, likely without Ca²⁺ binding function, as exemplified by the EF-hand-like of

PLCD4. Interestingly, the second EF-hand of Calnuc, which is the one overlapping with its GBA motif, is also considered a non-canonical EF-hand based on sequence divergence (66,67), and it only regulates G proteins when it is not bound to Ca²⁺ (25).

To assess protein-protein binding of PLC δ 4 isoforms to G proteins, lysates of HEK293T cells expressing myc-tagged PLC δ 4a and PLC δ 4b were used in pulldown assays with purified GST-Gai3. We found that PLC δ 4b binds Gai3 much more robustly than PLC δ 4a (**Fig. 9B**). A possible explanation for this difference is that the GBA motif is more accessible to the G protein in the shorter isoform (in which it is located near the C-terminal end) than in the longer one (in which it is sandwiched between the EF-hands and the catalytic TIM barrel). Regardless of the explanation for the difference in Gai3 binding among isoforms, these results suggest that PLC δ 4b might be a better regulator of G protein signaling in cells than PLC δ 4a.

PLC δ 4b but not PLC δ 4a promote G protein activation in cells- We used the BRET-based biosensor that directly monitors G protein activity described above (i.e., V-G β γ / GRK3ct-Nluc) to assess the function of PLC δ 4a and PLC δ 4b in cells. Myc-tagged constructs were expressed in HEK293T cells and BRET measured under steady-state conditions. Consistent with the protein-protein binding results, we found that expression of PLC δ 4b but not PLC δ 4a led to an increase in G protein activity as determined by BRET (**Fig. 9C**). This BRET enhancement was not reproduced by the PLC δ 4b FALA mutant (which is GEF-deficient, **Fig. 7D**), indicating that PLC δ 4b activates G proteins in mammalian cells via its GBA motif. Equivalent results were obtained using a construct consisting of PLCD4 aa 1 to 260 (**Fig. S7**), which is similar to PLC δ 4b and identical to the construct used in yeast-based experiments (**Figs. 2, 4**) and in *in vitro* protein binding experiments (**Fig. 3, 5**). These results further validate that PLCD4 exhibits consistent G protein regulatory activity across different experimental systems.

To further substantiate that PLC δ 4b activates G protein signaling in cells, we

investigated whether it regulates cAMP levels. For this, we fused full-length PLC δ 4b (and PLC δ 4a) to FKBP and tested the effect of recruiting the construct to the plasma membrane on forskolin-induced cAMP levels. We found that membrane recruitment of full-length PLC δ 4b led to a decrease in cAMP levels (**Fig. 10A, B**), much like the recruitment of its GBA motif (**Fig. 8E, F**). PLC δ 4b-mediated inhibition of cAMP was abolished upon introduction of the FALA mutation (**Fig. S8**), indicating that the observed response is GBA motif-dependent. The effect of full-length PLC δ 4b was more modest than that of its GBA motif alone (~12% vs. ~25% [**Fig. 8**], respectively), which might be explained by their relative levels of expression (**Fig. S8**). Based on our results from BRET-based G protein activity assays in **Fig. 9**, we predicted that PLC δ 4a would not inhibit forskolin-induced cAMP levels. Surprisingly, we found that, even though PLC δ 4a did not inhibit cAMP, it caused an *enhancement* of forskolin-induced cAMP levels (**Fig. 10A, B**). We reasoned that this enhancement of forskolin-induced cAMP levels was due to its catalytic PLC activity (which is absent in PLC δ 4b). This is not far-fetched because several AC isoforms are activated by increased Ca²⁺ levels or Ca²⁺-dependent mechanisms (59), a direct downstream effect of PLC activity. We found that the effect of PLC δ 4a on cAMP was indeed dependent on its PLC activity, as it was inhibited by the small molecule PLC inhibitor U73122 (**Fig. 10D**). On the other hand, U73122 had no effect on PLC δ 4b-mediated inhibition of forskolin-induced cAMP levels (**Fig. 10D**), which is consistent with the lack of a catalytic PLC domain in this isoform and supports the specificity of the effect of the inhibitor on PLC δ 4a. Taken together, these findings confirm that, instead of working as a phospholipase, PLC δ 4b is a non-receptor activator of heterotrimeric G proteins that functions via a defined GBA motif.

DISCUSSION

The main advances reported here are three: (i) the validation of a pipeline to identify and validate novel GBA motif-containing G protein regulators, (ii) the discovery of PLC δ 4b as a novel non-

receptor G protein activator with a GBA motif, and (iii) a systematic characterization of the biochemical, structural and signaling properties of all the GBA motifs described to date. Together, these findings advance our understanding of an understudied family of G protein regulators with important biological functions and establish a framework for their discovery and characterization.

The existence of a specific sequence that defines a group of non-receptor G protein activators (i.e., the GBA motif) is useful but insufficient information to reliably identify novel members of this family of G protein regulators based solely on sequence similarity. This contrasts with other families of G protein regulators like RGS GAPs, as many RGS proteins were discovered based on sequence similarity. This is because, while the GBA motif is a short sequence found in only a few proteins, RGS proteins are defined by a longer sequence (i.e., the RGS box, of about 120 aa) present in a larger number of proteins, thereby providing a more robust training set for bioinformatics searches. The findings reported here help to overcome the shortcomings of sequence-based searches of GBA regulators in two ways. The first is that increasing the number of GBA sequences that bind to G proteins will be helpful to improve the training set for future bioinformatics searches. Second, and probably more useful, is the pipeline of high and medium throughput assays to filter out false positives from sequence searches. Dozens or hundreds of GBA motif candidates can be rapidly assessed for the essential biochemical property of binding to G proteins by using peptide arrays, while yeast is used as a secondary system of facile genetic manipulation to test for G protein signaling activity. It is tempting to speculate about the implementation of a similar pipeline to discover new members of another family of G protein regulators such as GDIs that contain the so-called GoLoco motif. Much like the GBA motif family, these regulators are characterized by a short sequence (30-40 aa) present in a relatively small number of proteins, and the pipeline of assays described here is compatible with their characterization.

Related to the above, our comprehensive characterization of all the GBA motifs described to date in mammals (GIV, DAPLE, Calnuc, NUCB2 and, now, PLCD4) provides informative benchmarks and tools to assess the biochemical, structural and signaling properties of this family. Consistent with prior data obtained through the characterization of GIV's GBA motif by mutagenesis (23,29,31), the potency of different GBA motifs in triggering G protein activation in cells (**Fig. 8**) correlates with their affinity for G proteins *in vitro* (**Fig. 6**). Likewise, all share a common binding site on the G protein, although the newly identified GBA motif in PLC δ 4 has slightly different requirements to establish the protein-protein interaction. Collectively, it appears that the C-terminal region of the GBA motif downstream of PLC δ 4 F227, which diverges in sequence from other GBA motifs (**Fig. 6A**), might establish different protein-protein contacts, as indicated by the requirement of simultaneous mutation of F227 and L228 to alanine to completely disrupt binding and by the different impact of the Gai3 W258F mutation on PLC δ 4 binding compared to other GBA motifs. Elucidating such nuanced differences in the mode of G protein binding by GBA motifs will require atomic resolution structures.

The new GBA motif-containing protein we identified here, PLC δ 4, presents a number of interesting properties. The most striking one is that we have been able to validate the G protein regulatory activity for one of the two annotated isoforms, PLC δ 4b. While both PLC δ 4a and PLC δ 4b have been reported to be ubiquitously expressed in different tissues (61,62), PLC δ 4b appears to be a G protein regulator rather than a phospholipase. This is because this isoform is composed only of the PH domain and the EF-hands, lacking the catalytic TIM barrel required for lipase activity. Interestingly, it has been previously reported that the PH domain of PLC δ 4 in isolation, but not the full-length protein corresponding to PLC δ 4a, localizes to the plasma membrane via phosphoinositide binding—different manipulations that decrease PIP₂ levels lead to the release of PLC δ 4 PH domain from the plasma membrane into the cytosol (68). Because PLC δ 4b consists primarily of the PH domain with only a C-terminal extension containing the EF-

hands, it is likely that its subcellular localization and regulation by phosphoinositides recapitulates the properties of the PH domain in isolation. This raises the interesting possibility that the G protein regulatory activity of PLC δ 4b might be controlled by alterations in phosphoinositide levels. We have recently shown that the main mechanism by which GBA proteins regulate G protein signaling is by spatial proximity (36)—i.e., G protein activation occurs when the GBA motif is recruited to membranes in physical proximity to the target G protein, G α i, which is constitutively membrane-bound. Thus, if phosphoinositides regulate the localization of PLC δ 4b at the plasma membrane, they might also impact its G protein regulatory function. At a physiological level, it is not possible at this time to envision what function the GBA motif of PLC δ 4 might have. PLC δ 4 function has not been extensively characterized, but insights from mice in which the *PLCD4* gene has been knocked out indicate that it is required for male fertility (69,70). However, this function is likely exerted by the PLC δ 4a isoform and we have not observed G protein regulatory activity for this isoform. While we cannot rule out that the GBA motif of PLC δ 4a might become functional under specific circumstances that would make the GBA motif accessible to G proteins (as PLC enzymes are believed to undergo significant conformational and/or spatial rearrangements (71,72)), the only available evidence for G protein regulatory activity is for the PLC δ 4b isoform. Unfortunately, the function(s) of this isoform are still unknown. Further work will be required to elucidate the role of the GBA motif in the biological functions of PLC δ 4.

EXPERIMENTAL PROCEDURES

Reagents and Antibodies- Unless otherwise indicated all reagents were of analytical grade and obtained from Fisher Scientific or Sigma. Cell culture media and goat anti-rabbit and goat anti-mouse Alexa Fluor 680 antibodies were from Life Technologies. Fluorescein di- β -D-galactopyranoside was from Marker Gene Technologies, and the protein inhibitor mixture was from Sigma (catalog no. S8830). Leupeptin, pepstatin, and aprotinin were from Gold Biotechnology. All restriction endonucleases and

Escherichia coli strain BL21(DE3) were from Thermo Scientific. *E. coli* strain DH5 α was purchased from New England Biolabs. *Pfu* ultra DNA polymerase was purchased from Agilent. Rabbit antibodies raised against Gai3 (C-10), and G β (M-14) were from Santa Cruz Biotechnology (Santa Cruz, CA). Mouse monoclonal antibodies raised against α -tubulin (catalog no. T6074) and His-tag (catalog no. H1029) were from Sigma, and for HA tag (12CA5) were from Roche. Rabbit antibodies for ppERK (catalog no. 4370) and mouse antibodies for Myc (catalog no. 9B11) were from Cell Signaling. Rabbit polyclonal antibody for RFP was from Abcam (catalog no. 62341). Goat anti-rabbit and goat anti-mouse IRDye 800 F(ab')₂ were from Li-Cor Biosciences (Lincoln, NE).

Bioinformatics Searches and Analyses-

The search for proteins with a GBA motif in *H. sapiens* was performed in ScanSite 3 using the QuickMatrix Method (44). Briefly, we used the sequences of known GBA motifs of the proteins GIV, DAPLE, Calnuc, NUCB2, GBAS-1 and the synthetic peptides KB-752 and GSP (23,25-27,41,43) to design the pattern as [VLIM]-[T]-[VLIM]-x-[DE]-F-[VLIM] and generate the corresponding position-specific scoring matrix (PSSM) to search in the *H. sapiens* proteome in the NCBI Protein GenPept/RefSeq database.

Synthesis of Peptide Libraries and Gai3

Overlay- Libraries of immobilized peptides were produced by automatic SPOT synthesis on continuous cellulose membrane supports (Whatman 50 cellulose membranes) using Fmoc (fluoren-9-ylmethoxycarbonyl) chemistry with the AutoSpot-Robot ASS 222 (Intavis Bioanalytical Instruments AG) (73). Candidate GBA motif peptides were synthesized as 25-mer peptides in which the invariable phenylalanine (F) of the core of the GBA motif was in the central position. Individual peptide-cellulose complexes were solubilized and re-spotted on Celluspot® slides for subsequent probing. Slides were primed in binding buffer (4.3 mM Na₂HPO₄, 1.4 mM KH₂PO₄, pH 7.4, 137 mM NaCl, 2.7 mM KCl, 5mM MgCl₂, 1mM DTT, 30 μ M GDP, 1% (v:v) TX100) and blocked for 1 h in the same buffer supplemented with 5% (w:v) BSA. Slides were incubated for 2h at room temperature with rat His-Gai3 at 20 μ g/

ml (~0.5 μ M) in the same buffer. After 4 washes, slides were sequentially incubated with primary (rabbit anti-Gai3, 1: 250; 90 min) and secondary (goat anti-rabbit Alexa Fluor 680; 60 min) antibodies. As a control for non-specific detection of the peptide sequence by the primary and/or secondary antibody, a replicate peptide array slide was processed in parallel under identical conditions except that no His-Gai3 was added. Images were acquired and spot intensity quantified in an Odyssey infrared scanner (Li-Cor). Specific binding was quantified by subtracting the signal intensity for each spot in the slide without Gai3 incubation from the corresponding spot signals in the slide with Gai3 incubation, and normalized by dividing the resulting number by the intensity of the spot after Coomassie blue staining. Images were processed using the ImageJ software (NIH) and assembled for presentation using Photoshop and Illustrator (Adobe).

Plasmid Constructs- Cloning of GBA candidates into plasmids for expression in yeast, bacteria or, in some cases, mammalian cells was done using a ligation independent cloning (LIC) system (74). Briefly, sequences corresponding to the GBA candidates indicated in **Fig. S1** were amplified by PCR from plasmids obtained from OpenBiosystems, Addgene, GenScript or individual investigators. The amplified fragments contained extensions at the 5' and 3' ends that made them compatible with any of the following LIC plasmids: pLIC-YES2 (yeast, described in (27)), pLIC-GST (bacteria) and pLIC-myc (mammalian). pLIC-GST and pLIC-myc plasmids were kindly provided by J. Sondek (UNC-Chapel Hill) (75). pLIC-YES2-Ric-8A has been described before (27). Cloning of the pET28b-Gai3 and pGEX-Gai3 plasmids for the expression of rat His-Gai3 or GST-Gai3, respectively, in bacteria have been described previously (23). pET24a-Gai3 for the expression of human Gai3 has been described in (29). The GBA motif sequences of human GIV (aa 1671-1701), human DAPLE (aa 1661-1691), human Calnuc (aa 303-333), human NUCB2 (aa 304-334) or human PLCD4 (aa 213-243) inserted into the NruI/BamHI sites of the pmRFP-FKBP-pseudojanin plasmid (Addgene #37999), which results in the replacement of the pseudojanin sequence by the GBA motifs to generate constructs that encode for RFP-FKBP-

GBA fusions. Plasmids encoding the GBA motif of human PLCD4 (aa 213-243) or of human GIV (aa 1660-1701) fused to GST were generated by LIC using the pLIC-GST vector mentioned above. The pT7-5-Gai1 plasmid (which bears an internal hexahistidine tag in the b/c loop of Gai1) was kindly provided by H. Hamm (Vandebilt University) (76) and the pbb131 plasmid encoding N-myristoyl transferase (NMT) (77) was a gift from Maurine Linder (Cornell University). The plasmid encoding Lyn11-FRB has been described previously (36). pcDNA3.1-A₁R, pcDNA3.1-Venus(1-155)-Gγ₂ (VN-Gγ₂) and pcDNA3.1-Venus(155-239)-Gβ₁ (VC-Gβ₁) were kindly provided by N. Lambert (Augusta University, GA) (58). pcDNA3.1-masGRK3ct-Nluc and pcDNA3.1-Nluc-EPAC-VV (57) were a gift from K. Martemyanov (Scripps Research Institute, FL). The pcDNA3-Gai3 plasmids for the expression of rat Gai3 in mammalian cells and the pET28b-GIV (1660-1870) plasmid for the expression of His-GIV-CT in *E. coli* have been described previously (23,42). Plasmids for the expression of N-terminally tagged myc-PLCδ4a, myc-PLCδ4b and myc-PLCδ4 (1-260) were generated by PCR amplification from full-length PLCD4 and insertion into pLIC-myc. The primers for myc-PLCδ4b included a 3' extension to add the amino acids in its C-terminus that diverge from myc-PLCδ4a and precede the stop codon. Plasmids encoding FKBP-fusions of PLCδ4a and PLCδ4b were generated by PCR amplification and insertion into the NruI/BamHI sites of the pmRFP-FKBP-pseudojanin as described above for the isolated GBA motifs.

Yeast Strains and Manipulations- The previously described (48) *Saccharomyces cerevisiae* strain CY7967 [*MATa GPA1(1-41)-Gai3 far1Δ fus1p-HIS3 can1 stel4:trp1:LYS2 ste3Δ lys2 ura3 leu2 trp1 his3*] (kindly provided by James Broach, Penn State University) was used for all yeast experiments. The main features of this strain are that the only pheromone responsive GPCR (Ste3) is deleted, the endogenous Gα-subunit Gpa1 is replaced by a chimeric Gpa1(1-41)-human Gai3 (36-354) and the cell cycle arrest-inducing protein *far1* is deleted. In this strain, the pheromone response pathway can be upregulated by the ectopic expression of activators of human Gai3 and does not result in the cell cycle arrest

that occurs in the native pheromone response (47,48). Plasmid transformations were carried out using the lithium acetate method. CY7967 was first transformed with a centromeric plasmid (CEN TRP) encoding the LacZ gene under the control of the *FUS1* promoter, which is activated by the pheromone response pathway. The P_{FUS1}::LacZ-expressing strain was transformed with pLIC-YES2 plasmids (2μm, URA) encoding each one of the GBA candidate constructs or Ric-8A described in “Plasmid Constructs”. Double transformants were selected in synthetic defined (SD)-TRP-URA media. Individual colonies were inoculated into 3ml of SDGalactose-TRP-URA and incubated overnight at 30°C to induce the expression of the proteins of interest under the control of the galactose-inducible promoter of pLIC-YES2. This starting culture was used to inoculate 20 ml of SDGalactose-TRP-URA at 0.3 OD600. Exponentially growing cells (~0.7-0.8 OD600, 4-5 hours) were pelleted to prepare samples for subsequent assays (see “β-galactosidase Activity Assay” and “Yeast protein immunoblotting” below).

β-galactosidase Activity Assay- This assay was performed as described previously (78) with minor modifications. Pellets corresponding to 0.5 OD600 (in duplicates) were washed once with PBS + 0.1% (w:v) BSA and resuspended in 200 μl assay buffer (60 mM Na₂PO₄, 40 mM NaH₂PO₄, 10 mM KCl, 1 mM MgCl₂, 0.25% (v:v) β-mercaptoethanol, 0.01% (w:v) SDS, 10% (v:v) chloroform) and vortexed. 100 μl were transferred to 96-well plates and reactions started by the addition of 50 μl of the fluorogenic β-galactosidase substrate fluorescein di-β-D-galactopyranoside (FDG, 100 μM final). Fluorescence (Ex. 485 ± 10 nm/ Em. 528 ± 10 nm) was measured every 2 min for 90 min at 30°C in a Biotek H1 synergy plate reader. Enzymatic activity was calculated from the slope of fluorescence (arbitrary units) versus time (min). At least 3 independent experiments were measured for each condition and the results normalized (fold activation) to the activity in controls (strains carrying an empty pLIC-YES2 plasmid).

Yeast Protein Immunoblotting- This assay was performed as described previously (78,79) with minor modifications. Briefly, pellets

corresponding to 5 OD600 were washed once with PBS + 0.1% BSA and resuspended in 150 μ l of lysis buffer (10 mM Tris-HCl, pH 8.0, 10% (w/v) trichloroacetic acid (TCA), 25 mM NH_4OAc , 1 mM EDTA). 100 μ l of glass beads were added to each tube and vortexed at 4°C for 5 min. Lysates were separated from glass beads by poking a hole in the bottom of the tubes followed by centrifugation onto a new set of tubes. The process was repeated after the addition of 50 μ l of lysis buffer to wash the glass beads. Proteins were precipitated by centrifugation (10 min, 20,000 xg) and resuspended in 60 μ l of solubilization buffer (0.1 M Tris-HCl, pH 11.0, 3% SDS). Samples were boiled for 5 min, centrifuged (1 min, 20,000 xg) and 50 μ l of the supernatant transferred to new tubes containing 12.5 μ l of Laemmli sample buffer and boiled for 5 min. Proteins (~15-20 μ l per lane) were separated by SDS-PAGE, blocked in PBS supplemented with 5% BSA and analyzed by sequential incubation with primary and secondary antibodies. Primary antibodies were diluted as follows: ppERK (which recognizes yeast ppFus3): 1:2,500, myc: 1:1,000, and α -tubulin: 1:2,500. Secondary antibodies (Goat anti-mouse IRDye 800 F(ab')₂, Li-Cor Biosciences, and Goat anti-rabbit Alexa Fluor 680, LifeTechnologies) were used at 1:10,000. Images were acquired in an Odyssey infrared scanner (Li-Cor), processed using ImageJ software (NIH) and assembled for presentation using Photoshop and Illustrator (Adobe).

Protein Expression and Purification- All His-tagged and GST-tagged proteins were expressed in BL21(DE3) *E. coli* transformed with the corresponding plasmids by overnight induction at 23 °C with 1 mM isopropyl- β -D-1-thiogalactopyranoside (IPTG). Protein purification was carried out following previously described protocols (23,42). Briefly, bacteria pelleted from 1 L of culture were resuspended in 25 ml of buffer [50 mM NaH_2PO_4 , pH 7.4, 300 mM NaCl, 10 mM imidazole, 1% (v/v) Triton X-100 supplemented with protease inhibitor cocktail (Leupeptin 1 μ M, Pepstatin 2.5 μ M, Aprotinin 0.2 μ M, PMSF 1 mM)]. For rat or human Gai3, this buffer was supplemented with 25 μ M GDP and 5 mM MgCl_2 . After sonication (four cycles, with pulses lasting 20 s/cycle, and with 1 min interval between cycles to prevent heating), lysates were centrifuged at

12,000g for 20 min at 4 °C. The soluble fraction (supernatant) of the lysate was used for affinity purification on HisPur Cobalt or Glutathione Agarose resins (Pierce) and eluted with lysis buffer supplemented with 250 mM imidazole or with 50 mM Tris-HCl, pH 8, 100 mM NaCl, 30 mM reduced glutathione, respectively. GST-tagged proteins were dialyzed overnight at 4 °C against PBS. For rat or human His-Gai3, the buffer was exchanged for 20 mM Tris-HCl, pH 7.4, 20 mM NaCl, 1 mM MgCl_2 , 1 mM DTT, 10 μ M GDP, 5% (v/v) glycerol using a HiTrap Desalting column (GE Healthcare). Validation of the quality of the human Gai3 mutants used in **Fig. 7B** by trypsin protection assays was shown previously (29). All protein samples were aliquoted and stored at -80 °C. Myristoylated Gai1 was purified from BL21(DE3) *E. coli* bacteria co-expressing NMT as described above for His-Gai3, except that after the cobalt affinity purification step the eluate was subjected to ion exchange chromatography in a HiTrapQ HP column connected to an AKTA FPLC. Bovine retinal G $\beta\gamma$ (i.e., G $\beta_1\gamma_1$) was purified as described previously. Briefly, holotransducin was purified from rod outer segment membranes isolated from dark-adapted bovine retina as described previously (80). The G $\beta\gamma$ complex was separated from the α -subunit of transducin on a Hitrap Blue-sepharose column (GE healthcare) and further purified using anion exchange chromatography using a Hitrap-Q column as described earlier (81). Gai3-G $\beta\gamma$ heterotrimers were reconstituted by incubating rat His-Gai3 (20-25 μ M) with 2-fold molar excess of G $\beta\gamma$ overnight at 4°C in 20 mM Tris-HCl, pH 7.4, 20 mM NaCl, 1 mM MgCl_2 , 1 mM DTT, 10 μ M GDP, 5% (v/v) glycerol. Reconstituted heterotrimers were aliquoted and frozen at -80 °C.

In Vitro Protein Binding Assays- GST pulldown assays were carried out as described previously (25,42) with minor modifications. For **Fig. 3**, 20-25 μ g (~1.5-2 μ M final) of GST or GST-fused GBA candidates (described in “Plasmid Constructs”) were immobilized on glutathione agarose beads for 90 min at room temperature in PBS. Beads were washed twice with PBS, resuspended in 250 μ l of binding buffer (50 mM Tris-HCl, pH 7.4, 100 mM NaCl,

0.4% (v:v) NP-40, 10 mM MgCl₂, 5 mM EDTA, 2 mM DTT, 30 μM GDP) and incubated 4h at 4°C with constant tumbling in the presence of 5 μg (~0.5 μM final) of rat His-Gαi3. Beads were washed four times with 1 ml of wash buffer (4.3 mM Na₂HPO₄, 1.4 mM KH₂PO₄, pH 7.4, 137 mM NaCl, 2.7 mM KCl, 0.1% (v/v) Tween-20, 10 mM MgCl₂, 5 mM EDTA, 1 mM DTT and 30 μM GDP) and resin-bound proteins eluted with Laemmli sample buffer by incubation at 37°C for 10 min. The conditions for **Fig. 5** were the same except that His-Gαi3 was preincubated 3 hours at 30 °C in binding buffer supplemented with GDP (30 μM), GDP + AlF₄⁻ (30 μM AlCl₃, 10 mM NaF) or GTPγS (30 μM) for PLCD4 or 30 min at 4° C with GDP or GDP + AlF₄⁻ for NOC2L. The wash buffer was also supplemented with the same nucleotides. For **Fig. 7C** and **Fig. S3** 10 μg of the indicated GST-fused protein and 1 μg of the G protein were used. For **Fig. 9B** and **Fig. S2**, 30 μg of GST-Gαi3 or GST were used with the lysates of ~75% of a 10-cm dish of HEK293T cells transfected with myc-PLCδ4a, myc-PLCδ4b or other GBA proteins shown in **Fig. S2**. Proteins were separated by SDS-PAGE and transferred to PVDF membranes. After blocking with PBS supplemented with 5% non-fat milk, membranes were analyzed by Ponceau S staining (GST-fused proteins) or sequential incubation with primary and secondary antibodies. Primary anti-His or anti-myc antibodies were used at 1:2,500 or 1:1,000 dilution, respectively, and secondary antibodies (Goat anti-mouse IRDye 800 F(ab')₂, Li-Cor Biosciences) at 1:10,000. Immunoblot quantification was performed by infrared imaging following the manufacturer's protocols using an Odyssey imaging system (Li-Cor Biosciences). All Odyssey images were processed using the Image J software (NIH) and assembled for presentation using Photoshop and Illustrator (Adobe).

Peptide Synthesis- Peptides corresponding to the GBA motif of human GIV (residues 1671-1701, KTGSPGSEVVTLQQFLEESNKLTSTVQIKSSS), human DAPLE (residues 1662-1695, SASPSSEMTLEEFLEESNRSSPTHDTPTSCRD DL), human Calnuc (residues 303-333, NVDTNQDRLVTLEEFLLASTQRKEFGDTGEG W), human NUCB2 (residues 304-334, EVDTNKDRLVTLEEFLLKATEKKEFLEPDSWE

) human PLCD4 (residues 213-243 ESFSADGQKLTLLEFLDFLQEEQKERDCTSE) or human NOC2L (residues 5-35) were synthesized using the *in situ* neutralization protocol for Boc-Solid Phase Peptide Synthesis (Boc-SPPS) on a *p*-methylbenzhydrylamine (MBHA) resin (Novabiochem, 0.67 mmol/g, 100-200 mesh). Following chain elongation, 5,6-carboxyfluorescein was activated with HATU and DIEA (4, 4 and 8 equivalents regarding to the amount of peptidyl-resin, respectively) and coupled to the resin-bound peptides at 65 °C for 1 h to yield the carboxyfluorescein-labeled peptides. Peptides were cleaved from the resin using a solution of hydrofluoric acid (HF) containing 5% of anisole for 1 h at 0 °C. Next, the HF solution containing the peptides was removed under vacuum and the resulting residues crushed out with Et₂O and filtered. The collected solids were redissolved in a 50% CH₃CN/H₂O solution containing 0.1% of trifluoroacetic acid (TFA), frozen down and lyophilized. Crude peptides were purified by reverse phase-HPLC using a XBridge BEH C18 OBD prep column (130Å, 5 μm, 19 mm X 150 mm) at a flow of 20 mL/min using H₂O (0.1% TFA) and CH₃CN (0.1%TFA) as eluents. The identity and final purity (>97%) of the peptides was determined by analytical RP-HPLC and mass spectrometry (ESI-TOF).

Fluorescence Polarization (FP)-based Peptide Binding Assays- Fluorescence polarization measurements were carried out in 384-well plates (Black OptiPlate-384F, Perkin Elmer). His-Gαi3 proteins (0-32 μM) and fluorescently-labeled peptide (0.025 μM) were mixed at room temperature for 10 min in a final volume of 20 μl of binding buffer (50 mM Tris-HCl, pH 7.4, 100 mM NaCl, 0.4% (v:v) NP-40, 10 mM MgCl₂, 5 mM EDTA, 2 mM DTT) supplemented with 30 μM GDP. Fluorescence polarization (Ex. 485 ± 10 nm/ Em. 528 ± 10 nm) was measured every 2 min for 30 min at room temperature in a Biotek H1 synergy plate reader to ensure that the signals were stable in time. Fluorescence polarization at different times was averaged, normalized to maximal binding and fitted to a one site binding model to determine the equilibrium dissociation constant (K_d) using Prism (GraphPad).

Protein Structure Modeling and Docking- A model of Gαi3 bound to a portion of PLCD4's GBA motif [aa 219-231] was generated by protein-protein docking using ICM version 3.8-3 (Molsoft LLC., San Diego, CA). Independent human Gαi3 and PLCD4 GBA models were first generated by homology using the x-ray crystal structure of Gαi1 bound to the synthetic GBA peptide KB-752 (41) as a starting template (PDB: 1Y3A; chains B and F for Gαi3 and PLCD4, respectively). The side chain positions of the separate homology models were energetically minimized using a Monte Carlo based method after initial construction. Protein-protein docking of the PLCD4 peptide to Gαi3 was then conducted *in silico* using a two-stage fast Fourier transform method (ICM, Molsoft LLC.). The best scoring solution was chosen to represent the binding mode. Model images were generated with PyMOL Molecular Graphics System (Schrödinger, LLC.).

Steady-state GTPase Assay- This assay was performed as described previously (23,25,42). Briefly, His-Gαi3 (100 nM) was preincubated with different concentrations of GST-PLCD4 GBA (aa 213-243) for 15 min at 30°C in assay buffer [20 mM Na-HEPES, pH 8, 100 mM NaCl, 1 mM EDTA, 25 mM MgCl₂, 1 mM DTT, 0.05% (w/v) C₁₂E₁₀]. GTPase reactions were initiated at 30°C by adding an equal volume of assay buffer containing 1 μM [γ-³²P]GTP (~50 c.p.m./fmol). Duplicate aliquots (25 μl) were removed at 15 min and reactions stopped with 975 μl of ice-cold 5% (w/v) activated charcoal in 20 mM H₃PO₄, pH 3. Samples were then centrifuged for 10 min at 10,000 xg, and 500 μl of the resultant supernatant were scintillation counted to quantify released [³²P]Pi. Background [³²P]Pi detected at 15 min in the absence of G protein was subtracted from each reaction and data expressed as percentage of the Pi produced by His-Gαi3 in the absence of GST-PLCD4. Background counts were <5% of the counts detected in the presence of G proteins. Experiments with G protein heterotrimers were performed essentially the same way. Briefly, reconstituted Gαi3-Gβγ heterotrimers (50 nM/100 nM) were preincubated with His-GIV-CT (2 μM) or GST-PLCD4 GBA (6 μM) before starting the reactions by the addition of [γ-³²P]GTP as described above.

Bioluminescence Resonance Energy Transfer (BRET)-based G Protein Activation Assays-

BRET experiments were conducted as described previously (36,82). HEK293T cells (ATCC®, CRL-3216) were seeded on 6-well plates (~400,000 cells/well) coated with gelatin and after one day transfected using the calcium phosphate method with plasmids encoding the following constructs (DNA amounts in parenthesis): Venus(155-239)-Gβ₁ (VC-Gβ₁) (0.2 μg), Venus(1-155)-Gγ₂ (VN-Gγ₂) (0.2 μg), Gαi3 (1 μg) mas-GRK3ct-Nluc (0.2 μg) and Lyn11-FRB (3 μg) along with FKBP-GBA constructs (0.1 μg). Approximately 16-24 hours after transfection, cells were washed and gently scraped in warm PBS, centrifuged (5 min at 550g) and resuspended in Tyrode's solution (140 mM NaCl, 5 mM KCl, 1 mM MgCl₂, 1 mM CaCl₂, 0.37 mM NaH₂PO₄, 24 mM NaHCO₃, 10 mM HEPES pH 7.4, 0.1% glucose) at a concentration of ~10⁶ cells/ml. 25,000 cells were added to a white opaque 96-well plate (Opti-Plate, Perkin Elmer) and mixed with the nanoluciferase substrate Nano-Glo (Promega, final dilution 1:200) for 2 min before measuring luminescence. PolarSTAR plate reader (BMG Labtech) was used to measure luminescence signals at 460 ± 20 nm and 528 ± 10 nm at 28 °C and BRET calculated as the ratio between the emission intensity at 528 ± 10nm divided by the emission intensity at 460 ± 20nm. For the kinetic measurements shown in **Fig. 8C** and **Fig. S5**, BRET was determined every 0.24 s and results presented as increase in BRET after subtraction of the basal signal measured for 30 s before any stimulation [ΔBRET (baseline)]. For the steady-state BRET measurement shown in **Fig. 9C** and **Fig. S7**, results were presented as difference in BRET compared to the signal in cells not expressing PLCD4 [ΔBRET (no PLCD4)]. Protein samples from BRET experiments were prepared by centrifugation of HEK293T cells and resuspension in lysis buffer (20 mM HEPES, pH 7.2, 5 mM Mg(CH₃COO)₂, 125 mM K(CH₃COO), 0.4% Triton X-100, 1 mM DTT, and protease inhibitor mixture). After clearing by centrifugation at 14,000 × g at 4 °C for 10 min, protein concentration was determined by Bradford. Samples were supplemented with Laemmli sample buffer and boiled for 5 min. Proteins were separated by SDS-PAGE and transferred to PVDF membranes, which were sequentially incubated

with primary and secondary antibodies (goat anti-rabbit or anti-mouse coupled to Alexa Fluor 680 or IRDye 800, 1:10,000). The primary antibodies were used at the following dilutions: α -tubulin 1:2,500, myc 1:1,000, and RFP 1:1,000. Infrared imaging of immunoblots was performed according to the manufacturer's protocols using an Odyssey Infrared Imaging System (Li-Cor Biosciences). All Odyssey images were processed using ImageJ software (NIH) and assembled for presentation using Photoshop and Illustrator (Adobe).

Intracellular cAMP measurements- This assay was performed using the previously described BRET-based biosensor NLuc-EPAC-VV (51,57). HEK293T cells were seeded, transfected and harvested as described above ("*Bioluminescence Resonance Energy Transfer (BRET)-based G Protein Activation Assays*") except that the plasmids used were (quantities in parenthesis): NLuc-EPAC-VV (0.2 μ g), Lyn11-FRB (3 μ g) and FKBP-GBA constructs (0.1 μ g). A PolarSTAR plate reader (BMG Labtech) was used to measure luminescence signals at 460 ± 20

nm and 528 ± 10 nm at 28 °C every 4 s and BRET calculated as the ratio between the emission intensity at 528 ± 10 nm divided by the emission intensity at 460 ± 20 nm. Results were presented as the inverse of the BRET ratio after subtraction of the basal signal measured for 60 s before any stimulation (ΔBRET^{-1}). Forskolin (3 μ M) was added at 60 s and rapamycin (0.5 μ M) at 360 s as indicated in the figures. Protein sample immunoblotting was performed as described in "*Bioluminescence Resonance Energy Transfer (BRET)-based G Protein Activation Assays*".

Statistical Analyses- Each experiment was performed at least three times. The data shown are presented as means with *error bars* representing the S.E.M. or as one representative result of each biological replicate (as indicated in the figure legends). Statistical significance between various conditions was assessed with Student's *t* test. $p < 0.05$ was considered significant.

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CONFLICT OF INTEREST

The authors declare that they have no conflicts of interest with the contents of this article.

AUTHOR CONTRIBUTIONS: MM, SB, VDG, J-CP, AL, LG-N and MG-M conducted experiments. MM, SB and MG-M designed experiments. MM, SB, VDG and MG-M analyzed data. JBB-C and GSB provided critical reagents. MM and MG-M wrote the manuscript. MG-M conceived and supervised the project. All authors reviewed the results and approved the final version of the manuscript.

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FOOTNOTES

¹The abbreviations used are: GPCR, G protein-coupled receptor; GEF, guanine nucleotide exchange factor; GBA, Ga-Binding-and-Activating; GDI: Guanine nucleotide dissociation inhibitor; GAP: GTPase activating protein; Gα binding and activating; GIV: Gα Interacting, Vesicle-associated protein; DAPLE: Dvl-associating protein with a high frequency of leucines; BRET, Bioluminescence resonance energy transfer; AC, adenylyl cyclase; aa, amino acid.

FIGURE LEGENDS

FIGURE 1. Identification of putative GBA motifs by bioinformatics and peptide array screening. (A) Alignment of known GBA sequences of GIV, DAPLE, Calnuc, NUCB2, and GBAS-1 proteins and the synthetic peptides KB-752 and GSP. The background was shaded black or grey if the residue was identical or similar, respectively, in $\geq 50\%$ of the sequences. The invariable phenylalanine (F), required for G protein binding, is in red. The consensus sequence for the 7 amino acids of the core of the GBA motif is shown underneath the alignment (ψ , hydrophobic; x, any). (B) Discovery pipeline for the identification and validation of G protein activators with a GBA motif. Putative GBA motifs are identified by bioinformatics, tested for G protein binding in a GBA peptide array and the corresponding genes evaluated for G protein activation in yeast-based signaling assays. Subsequent steps include validation of direct binding of GBA motif-containing proteins to GDP-bound G proteins and assessment of GBA-dependency for the activation of signaling in yeast. (C) Peptide array screening for binding of GBA motif candidates to Gai3. 25-mer peptides corresponding to each one of the indicated GBA motif candidates (full sequences of positive hits shown in **Fig. S1**) were synthesized and immobilized on slides as one peptide per spot. The immobilized peptides were probed in batch with purified His-Gai3 (Gai3 +) and binding determined after sequential incubation with primary and secondary antibodies coupled to fluorescent probes. Control reactions in which the slides were incubated with primary and secondary antibodies but without probing with purified His-Gai3 (Gai3 -) were performed in parallel. A third slide was stained with Coomassie blue (C). The arbitrary units (a.u.) of Gai3 binding were determined by subtracting the signal in the “Gai3 -” condition from the signal in the “Gai3 +” condition, and normalizing to the Coomassie staining signal. Results are the average of 4 experiments (performed with 2 independent batches of peptide synthesis) and the error bars are the S.E.M. Two peptides corresponding to the characterized GBA proteins Calnuc and DAPLE were included as internal controls and the previously determined K_d 's for their interaction with Gai3 are indicated.

FIGURE 2. Assessment of GBA motif-containing gene candidates as G protein activators in “humanized” yeast. (A) Schematic of humanized yeast reporter system to determine G protein activation by candidate GBA proteins. The pheromone response pathway in *Saccharomyces cerevisiae* is regulated by GPCR-mediated activation of heterotrimeric G proteins, which induces Fus3 phosphorylation and transcriptional activation of the *FUS1* promoter. A genetically engineered strain lacking endogenous pheromone-sensitive GPCRs, expressing human Gai3 instead of the endogenous Gα Gpa1 and bearing a *LacZ* reporter under the control of the *FUS1* promoter was used to determine GPCR-independent G

protein activation by exogenously expressed GBA proteins or by Ric-8A (positive control). *FUS1* promoter activity was determined by β -galactosidase activity assays (panel B) and Fus3 activation was determined by phosphoERK (ppERK, which recognizes yeast ppFus3) immunoblotting (panel C). **(B)** The β -galactosidase activity of yeast cells expressing the cDNA of the indicated GBA candidates (see details for each construct in **Fig. S1**) was measured as described in “*Experimental procedures*” and normalized relative to the activity in cells expressing an empty vector. Results are the average of six independent experiments and the error bars represent the S.E.M. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ compared to vector using the Student’s *t* test. **(C)** Yeast cells expressing the cDNA of the indicated GBA candidates (see details for each construct in **Fig. S1**) were lysed and proteins analyzed by immunoblotting with the indicated antibodies as described in “*Experimental procedures*”. One representative experiment of at least 3 is shown. The red arrowheads indicate the expected position for the full length protein corresponding to each construct.

FIGURE 3. PLCD4, NOC2L and K1045 GBA candidates bind directly to Gai3. Approximately 20 μ g of the indicated purified GST-fused constructs (see details for each construct in **Fig. S1**) were immobilized on glutathione-agarose beads and incubated with 5 μ g of purified His-Gai3 in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input= 5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown.

FIGURE 4. PLCD4 and NOC2L activate G protein signaling in yeast via their GBA motif. **(A)** The β -galactosidase activity of yeast cells expressing wild-type (WT) or “FA” mutant versions of the indicated GBA candidates (PLCD4, *left*; NOC2L, *center*; K1045, *right*) was measured as described in “*Experimental procedures*” and normalized relative to the activity in cells expressing an empty vector. Results are the average of six independent experiments and the error bars represent the S.E.M. *** $P < 0.001$ compared to the vector control or #### $P < 0.001$ compared to WT using the Student’s *t* test. **(B)** Yeast cells expressing the same constructs as in (A) were lysed and proteins analyzed by immunoblotting with the indicated antibodies as described in “*Experimental procedures*”. One representative experiment of at least 3 is shown.

FIGURE 5. PLCD4 and NOC2L bind preferentially to GDP-bound Gai3 over GTP-bound Gai3. Approximately 20 μ g of purified GST-PLCD4 (*left*) or GST-NOC2L (*right*) (or GST, as negative control) immobilized on glutathione-agarose beads were incubated with 5 μ g of purified His-Gai3 preloaded with GDP (inactive), GDP + AlF₄[−] (mimics GTP-bound transition state) or GTP γ S (non-hydrolyzable GTP analog) as indicated. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input= 5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown.

FIGURE 6. The GBA motif of PLCD4 binds Gai3 with high affinity. **(A)** Alignment of GBA 31-mer peptide sequences of PLCD4, NOC2L, GIV, DAPLE, Calnuc and NUCB2 used in fluorescence polarization (FP) assays. The background was shaded black or grey if the residue was identical or similar, respectively, in $\geq 50\%$ of the sequences. The invariable phenylalanine (F), required for G protein binding, is in red. A bar diagram of the domains of PLCD4 and NOC2L is presented to show the position of the GBA motif in the context of the full proteins. **(B)** The indicated fluorescently-labeled GBA peptides were

incubated with increasing amounts of purified rat His-Gai3 in the presence of GDP and binding determined by FP. Data were normalized to maximal binding (except for NOC2L, for which data is presented as increase in millipolarization (mP) units) and fitted to a one-site binding model (solid lines) to determine the equilibrium dissociation constant (K_d). Results expressed as average \pm S.E.M. of $n=3$.

FIGURE 7. PLCD4 GBA motif binds to the SwII/ α 3 helix region of Gai3 and has GEF activity *in vitro*. (A) Different views of a structure homology model of the complex formed between Gai3 and the GBA motif of PLCD4 with details of the residues within or near the predicted protein-protein interface. *Left panel*, overview of the G protein (blue): GBA motif (orange) complex. PLCD4 GBA motif is predicted to bind to the groove formed between the SwII and α 3 helix. *Middle panels*, two representations of the same view of the Gai3:PLCD4 interface. On the view on the right, side chains of amino acids in the SwII/ α 3 helix pocket of Gai3 mutated in the experiments shown in (B) and displayed in red and the PLCD4 GBA motif is omitted for clarity. *Right panel*, view of conserved hydrophobic amino acid side chains in PLCD4 GBA motif predicted to make contact with hydrophobic residues in the SwII/ α 3 helix pocket of Gai3. (B) The effect of Gai3 mutations on PLCD4 binding is highly correlated with their effect on GIV and DAPLE binding. *Left panel*, Equilibrium dissociation constants (K_d) for the binding of purified human His-Gai3 mutants to the GBA motif of PLCD4 was determined from FP binding curves as described in “*Experimental procedures*”. Results are expressed as average \pm S.E.M. of $n=3-5$. Fold changes in the K_d compared to WT that were larger than 2 or smaller than 0.5 are indicated in red or green, respectively. *Middle panel*, heat map (green to red scale shown on top) of $\text{Log}_2(K_d^{\text{MUT}}/K_d^{\text{WT}})$ for each Gai3 mutant relative to WT comparing PLCD4 GBA binding to previously obtained data (29) for GIV or DAPLE GBA binding. *Right panels*, correlation plots of $\text{Log}_2(K_d^{\text{MUT}}/K_d^{\text{WT}})$ values for PLCD4 and GIV (top) or PLCD4 and DAPLE (bottom). (C) Mutation of hydrophobic residues in PLCD4 GBA motif impairs Gai3 binding. Approximately 10 μg of purified GST or GST-fused PLCD4 GBA motif (aa 213-243, WT or bearing the indicating mutations) were immobilized on glutathione-agarose beads and incubated with 1 μg of purified rat His-Gai3 in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input= 5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown. (D) PLCD4 GBA motif increases Gai3 (left) or Gai3-G $\beta\gamma$ heterotrimer (right) activity *in vitro*. The steady-state GTPase activity of purified rat His-Gai3 or His-Gai3-G $\beta\gamma$ was determined in the presence of GST-PLCD4 GBA motif WT (black) or FALA mutant (red) by measuring the production of [^{32}P]Pi at 15 min as described in “*Experimental procedures*”. The effect of His-GIV-CT (aa 1660-1870, containing the GBA motif) WT (black) or F1685A mutant (red) on the activity of the Gai3-G $\beta\gamma$ heterotrimer was also tested. Results are the average \pm S.E.M. of $n=3$. ** $P<0.01$ using the Student’s *t* test.

FIGURE 8. PLCD4 GBA motif activates G protein signaling in cells. (A) *Left*, Translocation of GBA sequences to membranes is controlled by CID using the FKBP-rapamycin-FRB system. FKBP-fused GBA is recruited to membranes upon rapamycin-induced binding of FKBP to the FRB domain that is fused to a membrane-targeting sequence (Lyn11). *Right*, G protein activation is determined by BRET. Dissociation of Gai3:G $\beta\gamma$ trimers upon G protein activation leads to the release of Venus-tagged G $\beta\gamma$ (V-G $\beta\gamma$), which binds to the C-terminal domain of GRK3 fused to nanoluciferase (GRK3ct-Nluc) and causes an increase in BRET signal. (B) Immunoblot of FKBP-fused GBA motifs (WT or FA mutants) of human GIV (aa 1671-1701), DAPLE (aa 1661-1691), Calnuc (aa 303-333), NUCB2 (aa 304-334) or PLCD4 (aa 213-243) expressed in HEK293T cells used for the experiments shown in (C). (C) Rapamycin-induced translocation of FKBP-fused WT GBA motifs (black) leads to G protein activation as determined by BRET, and FA mutations (red) abolish or diminish this response. HEK293T cells expressing all the required assay components and transfected with equal amounts of plasmids encoding the FKBP-GBA

fusions (0.1 μ g per well) were treated with rapamycin (0.5 μ M) at the indicated time point (arrow). Results are the average \pm S.E.M. (shown only at 5-s intervals for clarity) of $n=4$ expressed as increase in BRET after rapamycin stimulation (Δ BRET, baseline corrected by subtraction of the BRET values before stimulation). *Lower right panel*, scatter plot of Δ BRET 90 seconds after rapamycin addition for the indicated FKBP-GBA constructs. * $P<0.05$, ** $P<0.01$, *** $P<0.001$ compared to the respective WT using the Student's t test. The graph and immunoblot bands for FKBP-GBA PLCD4 have been reused in **Fig. S4** for comparison with other results. **(D)** *Left*, Translocation of GBA sequences to membranes is controlled by CID using the FKBP-rapamycin-FRB system and in (A). *Right*, cAMP levels are measured by BRET using the Nluc-EPAC-VV sensor. Gai3-GTP negatively regulates adenylyl cyclase (AC), thereby diminishing cAMP levels induced upon forskolin-mediated activation of AC. **(E)** Immunoblot of FKBP-fused GBA motifs of human GIV or PLCD4 expressed in HEK293T cells used for the experiments shown in (F). **(F)** Rapamycin-induced translocation of FKBP-GBA GIV or FKBP-GBA PLCD4 inhibits forskolin-induced cAMP levels. HEK293T cells expressing the cAMP sensor Nluc-EPAC-VV, Lyn11-FRB and the indicated FKBP-fused GBA constructs were treated at the indicated times (arrows) with forskolin alone (3 μ M, black), forskolin followed by rapamycin (0.5 μ M, red) or no compound (grey). One representative experiment is shown on the left graphs and a scatter plot of rapamycin-induced inhibition of forskolin-stimulated cAMP levels from $n=4$ is shown on the right. The scatter plot includes results of analogous experiments with cells expressing the α_{2A} adrenergic receptor and stimulated with brimonidine (5 μ M) instead of rapamycin for comparison. ** $P<0.01$, *** $P<0.001$ compared to the control cells (-) using the Student's t test.

FIGURE 9. PLC δ 4b binds Gai3 more efficiently than PLC δ 4a and activates G proteins in cells. **(A)** Bar diagrams of human PLC δ 4 isoforms and constructs. PLC δ 4a contains (from N to C terminus) a PH domain, two EF-hands, an EF-hand-like domain that overlaps with the GBA motif, a catalytic TIM barrel (with X- and Y-boxes) and a C2 domain. PLC δ 4b and PLC δ 4 (1-260) are truncated before the catalytic TIM barrel. *Inset*, partial sequence alignment of PLC δ 4a, PLC δ 4b and PLC δ 4 (1-260). The C-terminal sequence of PLC δ 4b differs from PLC δ 4a in the 14 aa preceding its stop codon, and the PLC δ 4 (1-260) construct used in previous experiments (Figs. 2 to 5) is very similar to PLC δ 4b. **(B)** Gai3 binding of two PLC δ 4 isoforms. Approximately 30 μ g of purified GST-Gai3 or GST immobilized on glutathione-agarose beads were incubated with lysates of HEK293T cells expressing either myc-PLC δ 4a (*left*) or myc-PLC δ 4b (*right*) in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Three different capture intensities (low, medium, and high) are shown for the myc IB. Input= 2.5% of the total amount of lysate added in each binding reaction. One representative experiment of at least three is shown. **(C)** PLC δ 4b, but not PLC δ 4a, activates G protein signaling in cells via its GBA motif. HEK293T cells were transfected with the indicated amounts of plasmids for the expression of myc-PLC δ 4a (WT) or myc-PLC δ 4b (WT or FALA mutants) and all the required components of the BRET-based G protein activity assay described in **Fig. 8A** (Gai3, V-G β γ and GRK3ct-Nluc). Results from $n=5$ are presented as the increase in BRET (Δ BRET) compared to cells not expressing PLC δ 4 in a box-and-whisker plot (median with boxes of 25-75%ile and error bars indicating the range. Cross marks indicate the average). An immunoblot (IB) of a representative experiment is shown below the graph. ** $P<0.01$ compared to the respective WT using the Student's t test.

FIGURE 10. PLC δ 4b inhibits forskolin-induced cAMP formation. **(A, B, C)** BRET-based cAMP measurements upon rapamycin-mediated recruitment of FKBP-PLC δ 4a or FKBP-PLC δ 4b were performed exactly as described in **Fig. 8D**. HEK293T cells expressing the cAMP sensor Nluc-EPAC-VV, Lyn11-FRB and the indicated FKBP-PLC δ 4 constructs (0.25 μ g DNA per well) were treated at the indicated times (arrows) with forskolin alone (3 μ M, black), forskolin followed by rapamycin (0.5 μ M,

red) or no compound (grey). One representative experiment is shown in (A) and a scatter plot of rapamycin-induced inhibition of forskolin-stimulated cAMP levels from $n=3$ is shown in (B). $**P<0.01$ compared to the control cells (-) using the Student's t test. An immunoblot of cell lysates from one representative experiment is shown in (C). **(D)** BRET-based cAMP measurements upon rapamycin-mediated recruitment of FKBP-PLC δ 4a or FKBP-PLC δ 4b were performed and quantified as described in (A, B, C) in the presence of the PLC inhibitor U73122 (+) (10 μ M, 60 min preincubation) or DMSO (-). Results are the average \pm S.E.M. of $n=4$. $**P<0.01$, ns= not significant compared to control cells pretreated with DMSO (-) using the Student's t test.

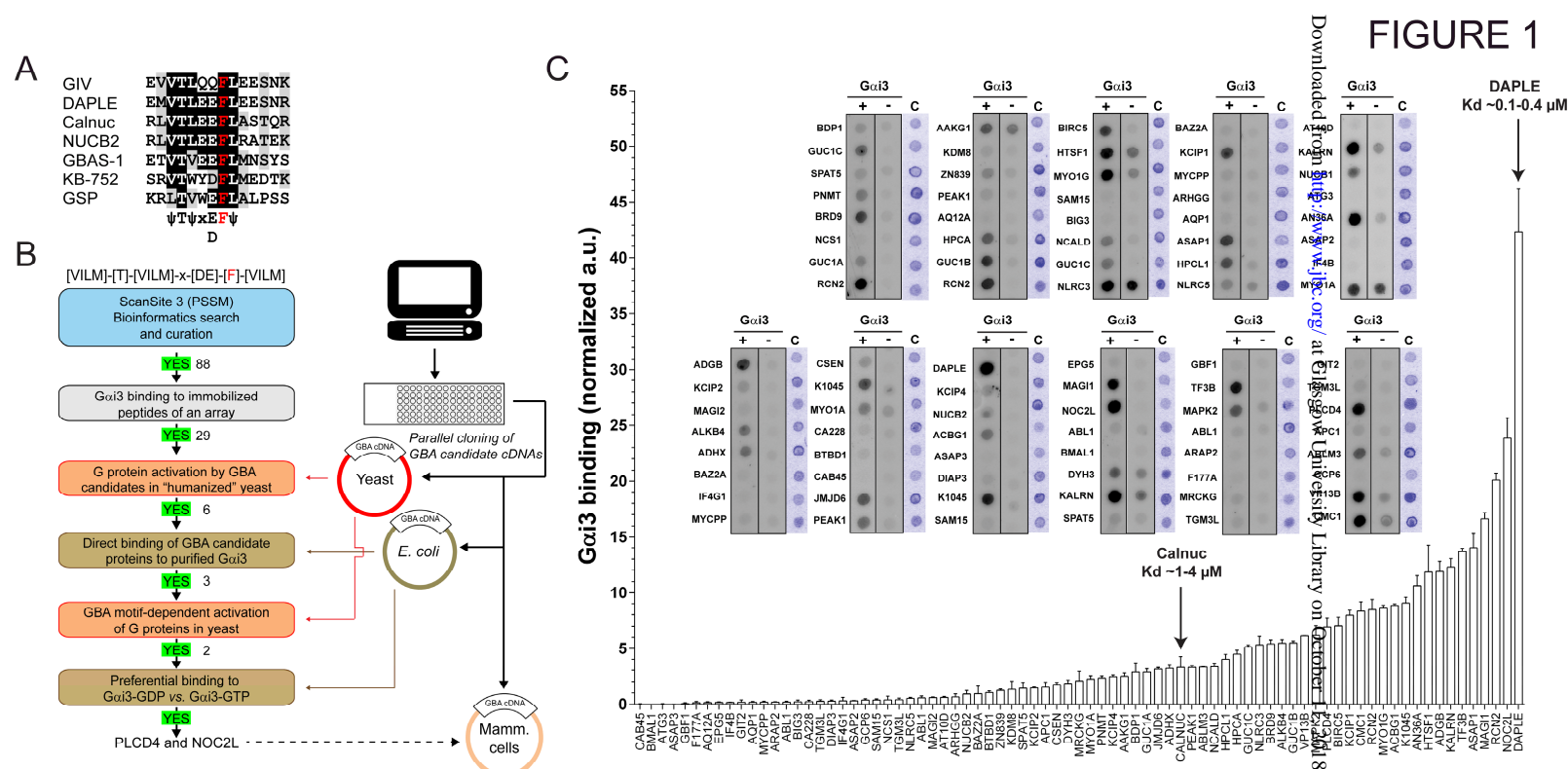


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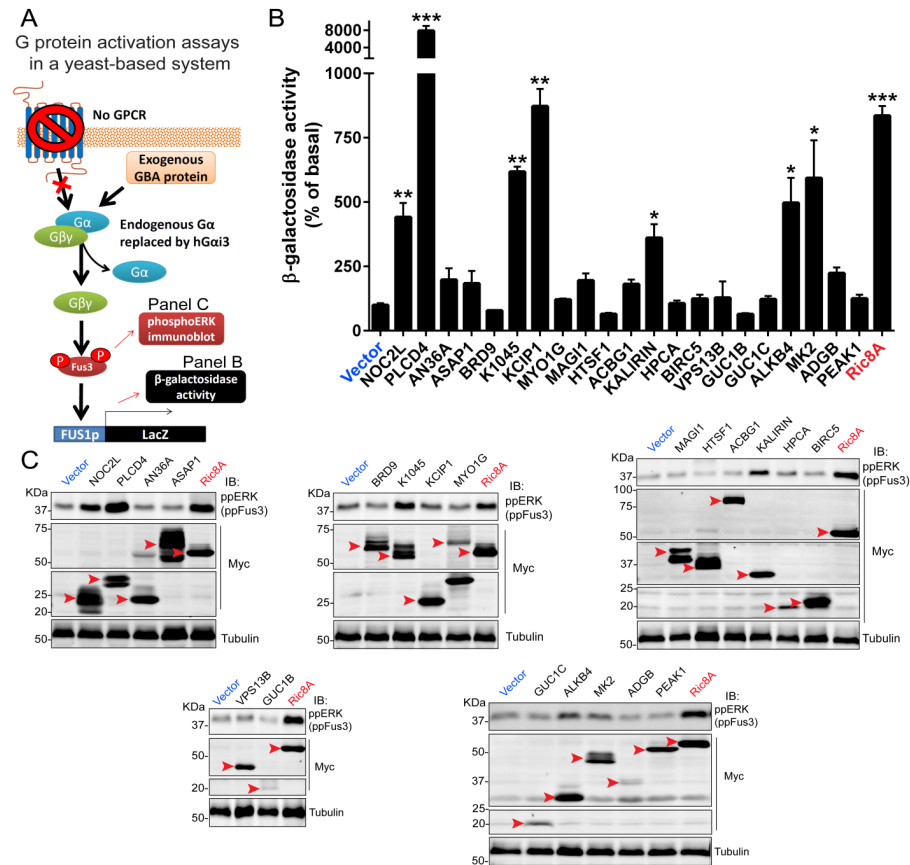


FIGURE 2. Assessment of GBA motif-containing gene candidates as G protein activators in “humanized” yeast. (A) Schematic of humanized yeast reporter system to determine G protein activation by candidate GBA proteins. The pheromone response pathway in *Saccharomyces cerevisiae* is regulated by GPCR-mediated activation of heterotrimeric G proteins, which induces Fus3 phosphorylation and transcriptional activation of the *FUS1* promoter. A genetically engineered strain lacking endogenous pheromone-sensitive GPCRs, expressing human Gai3 instead of the endogenous Gα Gpa1 and bearing a *LacZ* reporter under the control of the *FUS1* promoter was used to determine GPCR-independent G protein activation by exogenously expressed GBA proteins or by Ric-8A (positive control). *FUS1* promoter activity was determined by β-galactosidase activity assays (panel B) and Fus3 activation was determined by phosphoERK (ppERK, which recognizes yeast ppFus3) immunoblotting (panel C). (B) The β-galactosidase activity of yeast cells expressing the cDNA of the indicated GBA candidates (see details for each construct in Fig. S1) was measured as described in “Experimental procedures” and normalized relative to the activity in cells expressing an empty vector. Results are the average of six independent experiments and the error bars represent the S.E.M. *P < 0.05, **P < 0.01, ***P < 0.001 compared to vector using the Student’s *t* test. (C) Yeast cells expressing the cDNA of the indicated GBA candidates (see details for each construct in Fig. S1) were lysed and proteins analyzed by immunoblotting with the indicated antibodies as described in “Experimental procedures”. One representative experiment of at least 3 is shown. The red arrowheads indicate the expected position for the full length protein corresponding to each construct.

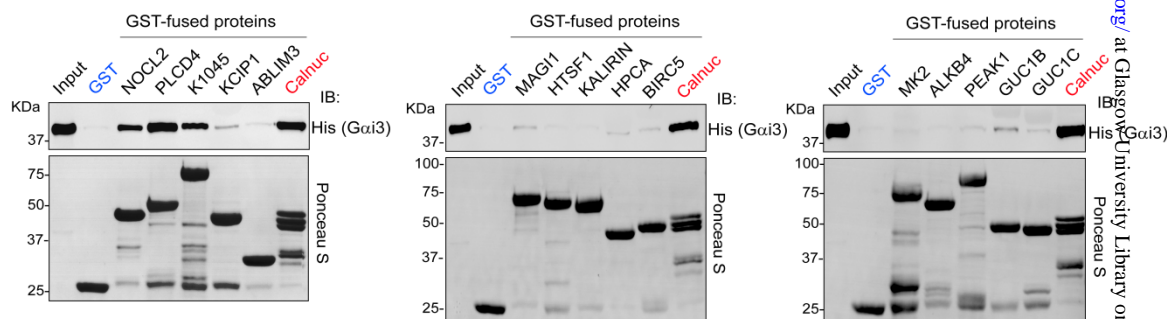


FIGURE 3. PLCD4, NOC2L and K1045 GBA candidates bind directly to Gai3. Approximately 20 μ g of the indicated purified GST-fused constructs (see details for each construct in **Fig. S1**) were immobilized on glutathione-agarose beads and incubated with 5 μ g of purified His-Gai3 in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input= 5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown.

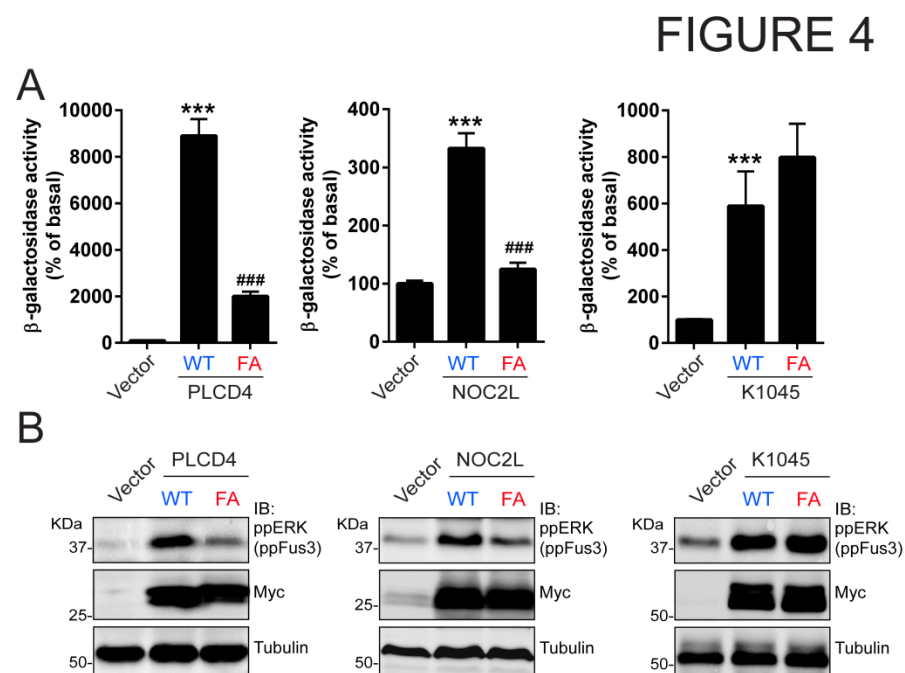


FIGURE 4. PLCD4 and NOC2L activate G protein signaling in yeast via their GBA motif. (A) The β -galactosidase activity of yeast cells expressing wild-type (WT) or “FA” mutant versions of the indicated GBA candidates (PLCD4, *left*; NOC2L, *center*; K1045, *right*) was measured as described in “*Experimental procedures*” and normalized relative to the activity in cells expressing an empty vector. Results are the average of six independent experiments and the error bars represent the S.E.M. *** $P < 0.001$ compared to the vector control or ### $P < 0.001$ compared to WT using the Student’s *t* test. (B) Yeast cells expressing the same constructs as in (A) were lysed and proteins analyzed by immunoblotting with the indicated antibodies as described in “*Experimental procedures*”. One representative experiment of at least 3 is shown.

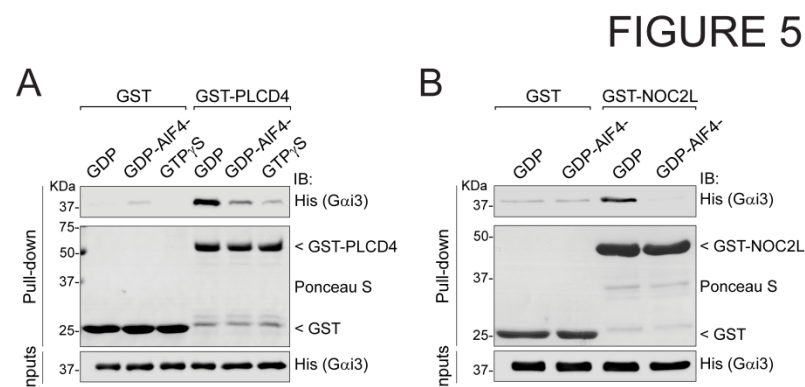


FIGURE 5. PLCD4 and NOC2L bind preferentially to GDP-bound Gai3 over GTP-bound Gai3. Approximately 20 μ g of purified GST-PLCD4 (*left*) or GST-NOC2L (*right*) (or GST, as negative control) immobilized on glutathione-agarose beads were incubated with 5 μ g of purified His-Gai3 preloaded with GDP (inactive), GDP + AIF₄⁻ (mimics GTP-bound transition state) or GTPγS (non-hydrolyzable GTP analog) as indicated. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input= 5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown.

FIGURE 6

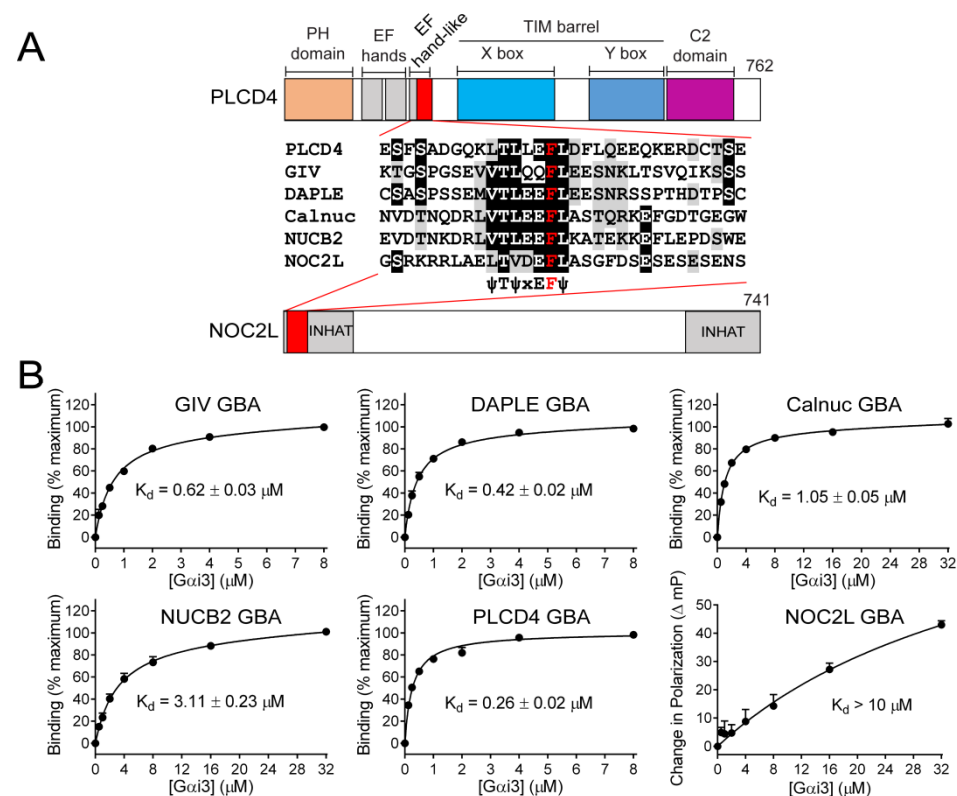


FIGURE 6. The GBA motif of PLCD4 binds Gαi3 with high affinity. (A) Alignment of GBA 31-mer peptide sequences of PLCD4, NOC2L, GIV, DAPLE, Calnuc and NUCB2 used in fluorescence polarization (FP) assays. The background was shaded black or grey if the residue was identical or similar, respectively, in $\geq 50\%$ of the sequences. The invariable phenylalanine (F), required for G protein binding, is in red. A bar diagram of the domains of PLCD4 and NOC2L is presented to show the position of the GBA motif in the context of the full proteins. (B) The indicated fluorescently-labeled GBA peptides were incubated with increasing amounts of purified rat His-Gαi3 in the presence of GDP and binding determined by FP. Data were normalized to maximal binding (except for NOC2L, for which data is presented as increase in millipolarization (mP) units) and fitted to a one-site binding model (solid lines) to determine the equilibrium dissociation constant (K_d). Results expressed as average \pm S.E.M. of $n=3$.

FIGURE 7

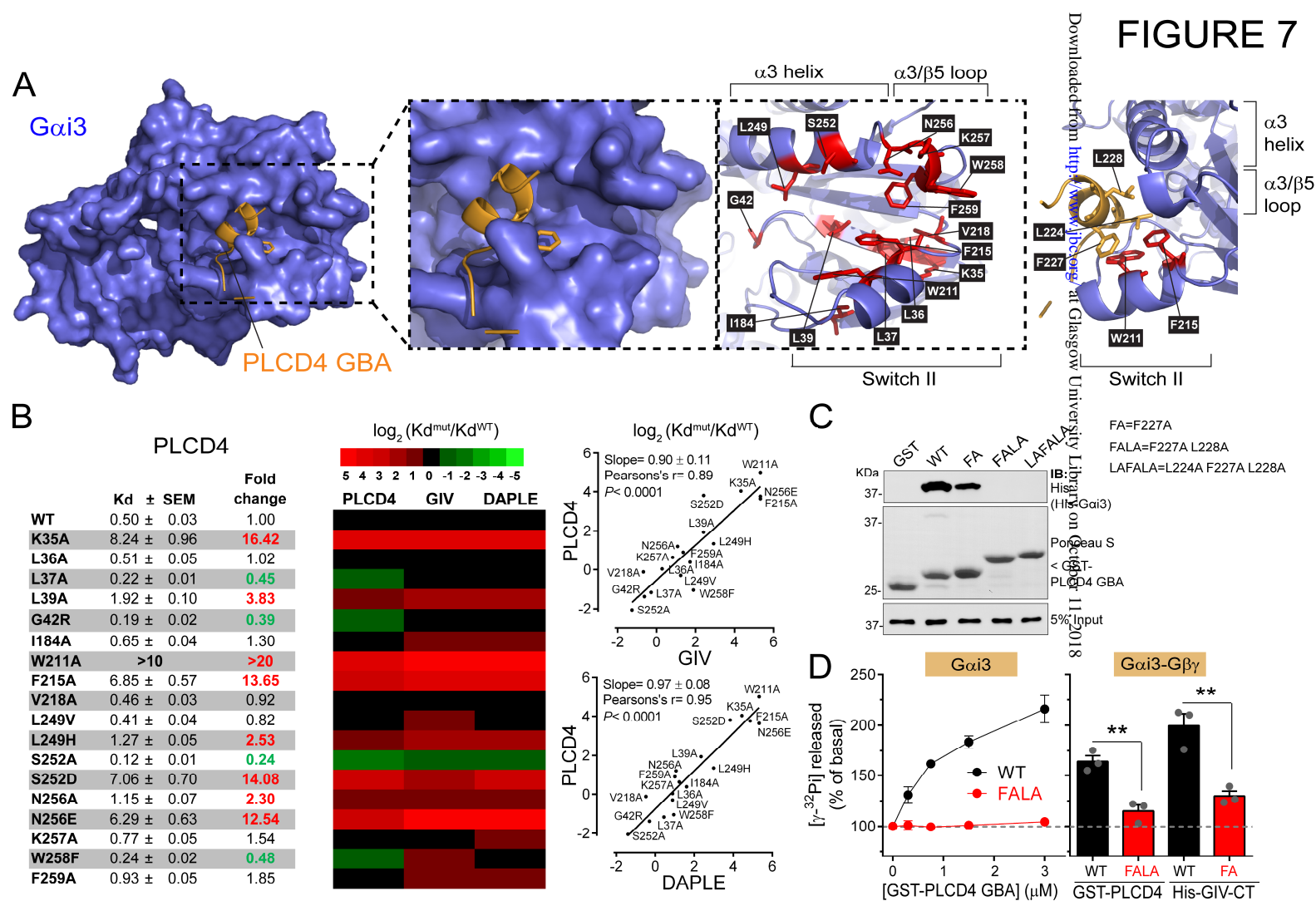


FIGURE 7. PLCD4 GBA motif binds to the SwII/ α 3 helix region of Gai3 and has GEF activity *in vitro*. (A) Different views of a structure homology model of the complex formed between Gai3 and the GBA motif of PLCD4 with details of the residues within or near the predicted protein-protein interface. *Left panel*, overview of the G protein (blue): GBA motif (orange) complex. PLCD4 GBA motif is predicted to bind to the groove formed between the SwII and α 3 helix. *Middle panels*, two representations of the same view of the Gai3:PLCD4 interface. On the view on the right, side chains of amino acids in the SwII/ α 3 helix pocket of Gai3 mutated in the experiments shown in (B) and displayed in red and the PLCD4 GBA motif is omitted for clarity. *Right panel*, view of conserved hydrophobic amino acid side chains in PLCD4 GBA motif predicted to make contact with hydrophobic residues in the SwII/ α 3 helix pocket of Gai3. (B) The effect of Gai3 mutations on PLCD4 binding is highly correlated with their effect on GIV and DAPLE binding. *Left panel*, Equilibrium dissociation constants (K_d) for the binding of purified human His-Gai3 mutants to the GBA motif of PLCD4 was determined from FP binding curves as described in “*Experimental procedures*”. Results are expressed as average \pm S.E.M. of $n=3-5$. Fold changes in the K_d compared to WT that were larger than 2 or smaller than 0.5 are indicated in red or green, respectively. *Middle panel*, heat map (green to red scale shown on top) of $\text{Log}_2(K_d^{\text{MUT}}/K_d^{\text{WT}})$ for each Gai3 mutant relative to WT comparing PLCD4 GBA binding to previously obtained data (29) for GIV or DAPLE GBA binding. *Right panels*, correlation plots of $\text{Log}_2(K_d^{\text{MUT}}/K_d^{\text{WT}})$ values for PLCD4 and GIV (top) or PLCD4 and DAPLE (bottom). (C) Mutation of hydrophobic residues in PLCD4 GBA motif impairs Gai3 binding. Approximately 10 μg of purified GST or GST-fused PLCD4 GBA motif (aa 213-243, WT or bearing the indicating mutations) were immobilized on glutathione-agarose beads and incubated with 1 μg of purified rat His-Gai3 in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Input=5% of the total amount of His-Gai3 added in each binding reaction. One representative experiment of at least three is shown. (D) PLCD4 GBA motif increases Gai3 (left) or Gai3-G $\beta\gamma$ heterotrimer (right) activity *in vitro*. The steady-state GTPase activity of purified rat His-Gai3 or His-Gai3-G $\beta\gamma$ was determined in the presence of GST-PLCD4 GBA motif WT (black) or FALA mutant (red) by measuring the production of [^{32}P]Pi at 15 min as described in “*Experimental procedures*”. The effect of His-GIV-CT (aa 1660-1870, containing the GBA motif) WT (black) or F1685A mutant (red) on the activity of the Gai3-G $\beta\gamma$ heterotrimer was also tested. Results are the average \pm S.E.M. of $n=3$. ** $P<0.01$ using the Student’s t test.

FIGURE 8

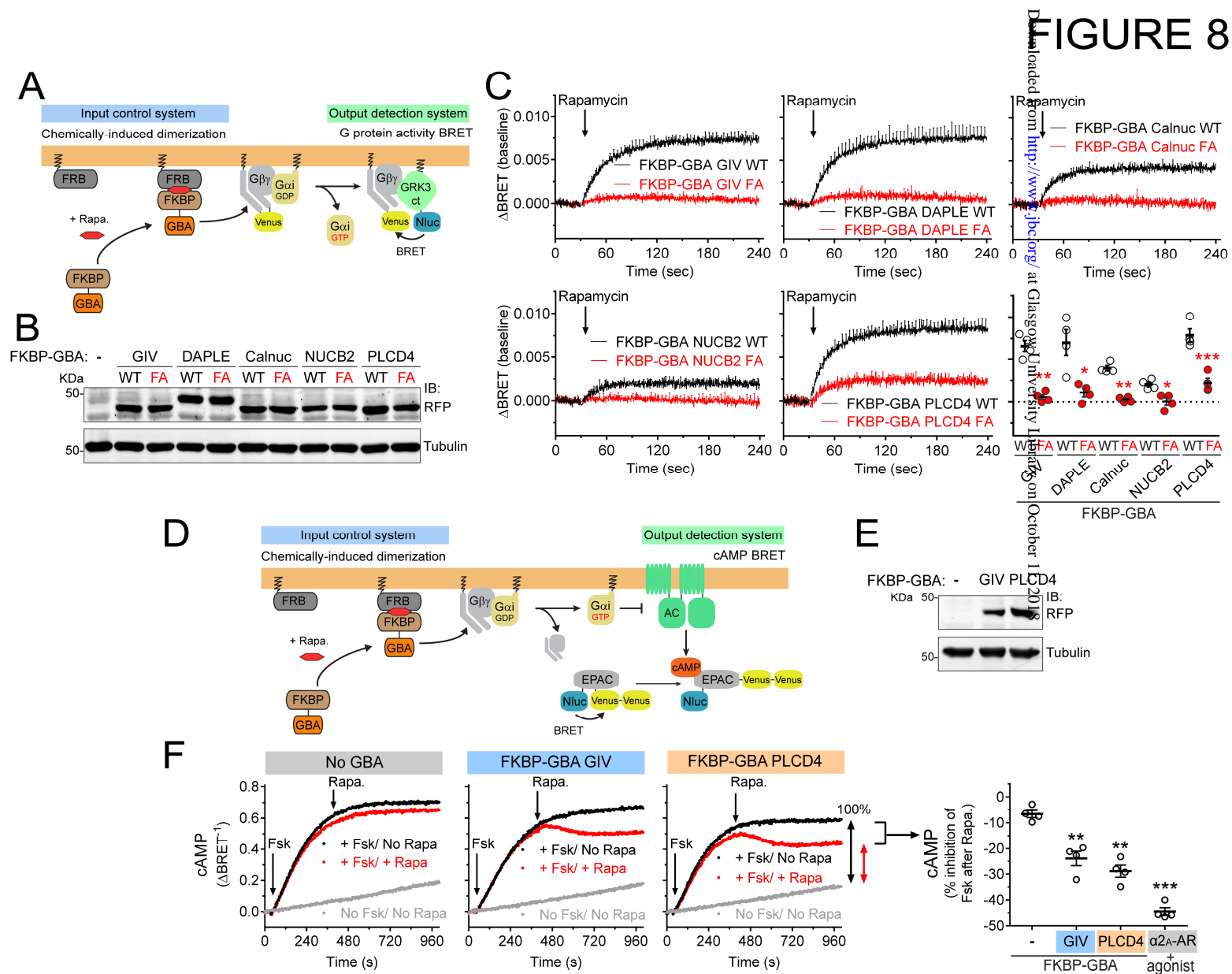


FIGURE 8. PLCD4 GBA motif activates G protein signaling in cells. (A) *Left*, Translocation of GBA sequences to membranes is controlled by CID using the FKBP-rapamycin-FRB system. FKBP-fused GBA is recruited to membranes upon rapamycin-induced binding of FKBP to the FRB domain that is fused to a membrane-targeting sequence (Lyn11). *Right*, G protein activation is determined by BRET. Dissociation of Gai3:Gβγ trimers upon G protein activation leads to the release of Venus-tagged Gβγ (V-Gβγ), which binds to the C-terminal domain of GRK3 fused to nanoluciferase (GRK3ct-Nluc) and causes an increase in BRET signal. (B) Immunoblot of FKBP-fused GBA motifs (WT or FA mutants) of human GIV (aa 1671-1701), DAPLE (aa 1661-1691), Calnuc (aa 303-333), NUCB2 (aa 304-334) or PLCD4 (aa 213-243) expressed in HEK293T cells used for the experiments shown in (C). (C) Rapamycin-induced translocation of FKBP-fused WT GBA motif (black) leads to G protein activation as determined by BRET, and FA mutations (red) abolish or diminish this response. HEK293T cells expressing all the required assay components and transfected with equal amounts of plasmids encoding the FKBP-GBA fusions (0.1 μg per well) were treated with rapamycin (0.5 μM) at the indicated time point (arrow). Results are the average ± S.E.M. (shown only at 5-s intervals for clarity) of n=4 expressed as increase in BRET after rapamycin stimulation (ΔBRET, baseline corrected by subtraction of the BRET values before stimulation). *Lower right panel*, scatter plot of ΔBRET 90 seconds after rapamycin addition for the indicated FKBP-GBA constructs. *P<0.05, **P<0.01, ***P<0.001 compared to the respective WT using the Student's *t* test. The graph and immunoblot bands for FKBP-GBA PLCD4 have been reused in Fig. S4 for comparison with other results. (D) *Left*, Translocation of GBA sequences to membranes is controlled by CID using the FKBP-rapamycin-FRB system and in (A). *Right*, cAMP levels are measured by BRET using the Nluc-EPAC-VV sensor. Gai3-GTP negatively regulates adenylyl cyclase (AC), thereby diminishing cAMP levels induced upon forskolin-mediated activation of AC. (E) Immunoblot of FKBP-fused GBA motifs of human GIV or PLCD4 expressed in HEK293T cells used for the experiments shown in (F). (F) Rapamycin-induced translocation of FKBP-GBA GIV or FKBP-GBA PLCD4 inhibits forskolin-induced cAMP levels. HEK293T cells expressing the cAMP sensor Nluc-EPAC-VV, Lyn11-FRB and the indicated FKBP-fused GBA constructs were treated at the indicated times (arrows) with forskolin alone (3 μM, black), forskolin followed by rapamycin (0.5 μM, red) or no compound (grey). One representative experiment is shown on the left graphs and a scatter plot of rapamycin-induced inhibition of forskolin-stimulated cAMP levels from n=4 is shown on the right. The scatter plot includes results of analogous experiments with cells expressing the α_{2A} adrenergic receptor and stimulated with brimonidine (5 μM) instead of rapamycin for comparison. **P<0.01, ***P<0.001 compared to the control cells (-) using the Student's *t* test.

FIGURE 9

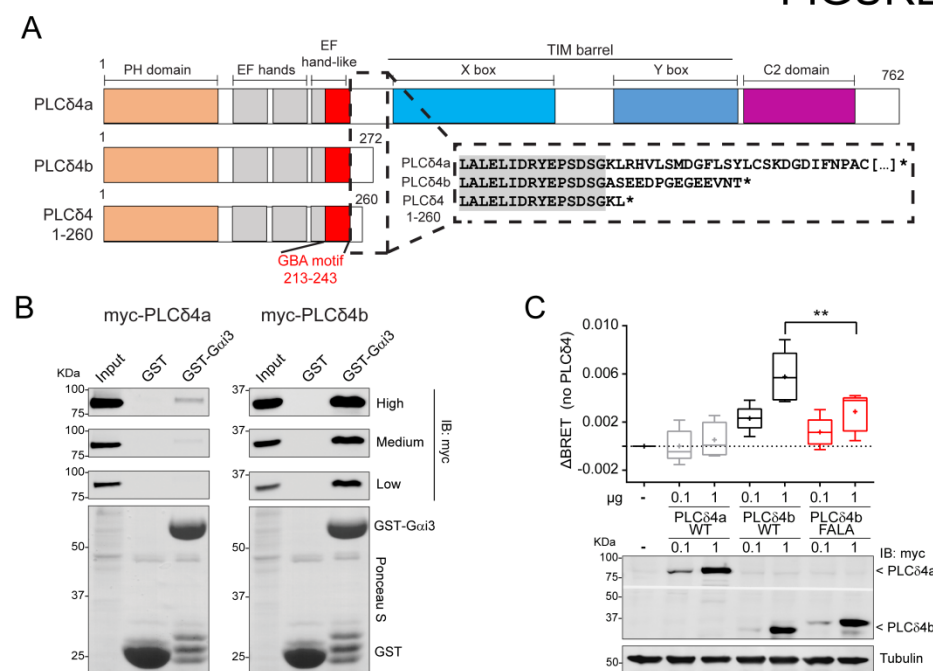


FIGURE 9. PLCδ4b binds Gai3 more efficiently than PLCδ4a and activates G proteins in cells. (A) Bar diagrams of human PLCδ4 isoforms and constructs. PLCδ4a contains (from N to C terminus) a PH domain, two EF-hands, an EF-hand-like domain that overlaps with the GBA motif, a catalytic TIM barrel (with X- and Y-boxes) and a C2 domain. PLCδ4b and PLCδ4 (1-260) are truncated before the catalytic TIM barrel. *Inset*, partial sequence alignment of PLCδ4a, PLCδ4b and PLCδ4 (1-260). The C-terminal sequence of PLCδ4b differs from PLCδ4a in the 14 aa preceding its stop codon, and the PLCδ4 (1-260) construct used in previous experiments (Figs. 2 to 5) is very similar to PLCδ4b. **(B)** Gai3 binding of two PLCδ4 isoforms. Approximately 30 μg of purified GST-Gai3 or GST immobilized on glutathione-agarose beads were incubated with lysates of HEK293T cells expressing either myc-PLCδ4a (*left*) or myc-PLCδ4b (*right*) in the presence of GDP. Resin-bound proteins were eluted, separated by SDS-PAGE, and analyzed by Ponceau S-staining and immunoblotting (IB) with the indicated antibodies. Three different capture intensities (low, medium, and high) are shown for the myc IB. Input = 2.5% of the total amount of lysate added in each binding reaction. One representative experiment of at least three is shown. **(C)** PLCδ4b, but not PLCδ4a, activates G protein signaling in cells via its GBA motif. HEK293T cells were transfected with the indicated amounts of plasmids for the expression of myc-PLCδ4a (WT) or myc-PLCδ4b (WT or FALA mutants) and all the required components of the BRET-based G protein activity assay described in Fig. 8A (Gai3, V-Gβγ and GRK3ct-Nluc). Results from n=5 are presented as the increase in BRET (ΔBRET) compared to cells not expressing PLCδ4 in a box-and-whisker plot (median with boxes of 25-75%ile and error bars indicating the range). Cross marks indicate the average. An immunoblot (IB) of a representative experiment is shown below the graph. **P<0.01 compared to the respective WT using the Student's *t* test.

FIGURE 10

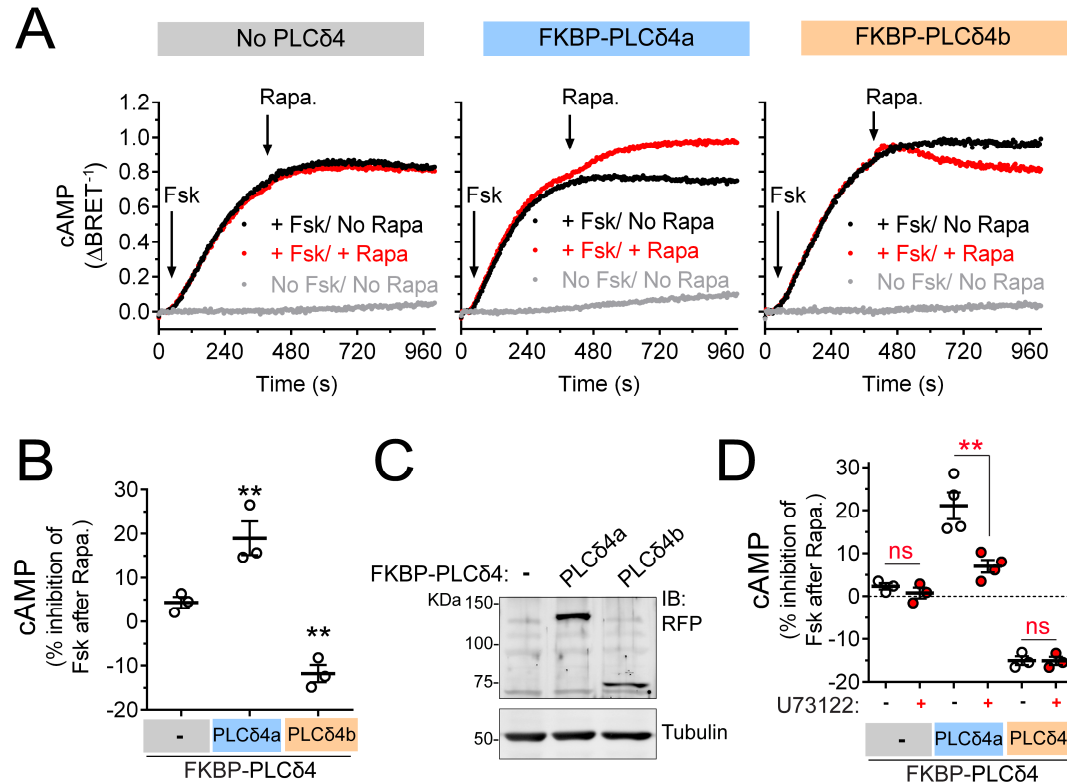


FIGURE 10. PLCδ4b inhibits forskolin-induced cAMP formation. (A, B, C) BRET-based cAMP measurements upon rapamycin-mediated recruitment of FKBP-PLCδ4a or FKBP-PLCδ4b were performed exactly as described in **Fig. 8D**. HEK293T cells expressing the cAMP sensor Nluc-EPAC-VV, Lyn11-FRB and the indicated FKBP-PLCδ4 constructs (0.25 μg DNA per well) were treated at the indicated times (arrows) with forskolin alone (3 μM, black), forskolin followed by rapamycin (0.5 μM, red) or no compound (grey). One representative experiment is shown in (A) and a scatter plot of rapamycin-induced inhibition of forskolin-stimulated cAMP levels from n=3 is shown in (B). **P<0.01, ***P<0.001 compared to the control cells (-) using the Student's *t* test. An immunoblot of cell lysates from one representative experiment is shown in (C). (D) BRET-based cAMP measurements upon rapamycin-mediated recruitment of FKBP-PLCδ4a or FKBP-PLCδ4b were performed and quantified as described in (A, B, C) in the presence of the PLC inhibitor U73122 (+) (10 μM, 60 min preincubation) or DMSO (-). Results are the average ± S.E.M. of n=4. **P<0.01, ns= not significant compared to control cells pretreated with DMSO (-) using the Student's *t* test.

A biochemical and genetic discovery pipeline identifies PLC δ 4b as a non-receptor activator of heterotrimeric G proteins

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